## 1 APPEARANCES 2 PANEL MEMBERS: 3 Arthur Fong, Ph.D., IBM Corporation, Co-Chairperson 4 Kelly Moran, Ph.D., TDC Environmental, LLC, Co-Chairperson 5 Caroline "Cal" Baier-Anderson, Ph.D., U.S. Environmental 6 Protection Agency 7 Ann Blake, Ph.D., Environmental and Public Health Consulting 8 Michael Caringello, S.C. Johnson & Son 9 Bill Carroll, Ph.D., Occident Chemical Corporation 10 (via teleconference) Ken Geiser, Ph.D., University of Massachusetts-Lowell 11 Helen Holder, Hewlett-Packard Company 12 13 Tim Malloy, J.D., University of California, Los Angeles, School of Law 14 Julia Quint, Ph.D., California Department of Public 15 Health(Retired) 16 Megan Schwarzman, M.D., M.P.H., University of California, Berkeley 17 Rebecca Sutton, Ph.D., San Francisco Estuary Institute 18 Don Versteeg, Ph.D., Procter & Gamble Company 19 Ken Zarker, Washington State Department of Ecology 20 21 STAFF: 22 Debbie Raphael, Director 23 Meredith Williams, Ph.D., Deputy Director, Safer Products and Workplaces Program 2.4

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## APPEARANCES CONTINUED STAFF: Andre Algazi, Senior Environmental Scientist Bob Boughton, Senior Hazardous Substances Engineer Lynn Goldman, Staff Counsel Radhika, Majhail, Public Participation Specialist Karl Palmer, Branch Chief Hortensia Muniz, Senior Hazardous Substances Engineer ALSO PRESENT: Greg Gorder, Technology Sciences Group Will Lorenz, General Coatings Miriam Rotkin-Ellman, Natural Resources Defense Counsel

## PROCEEDINGS

PUBLIC PARTICIPATION SPECIALIST MAJHAIL: Okay.

Good morning, everybody. How are you all today?

You look rested.

(Laughter.)

PUBLIC PARTICIPATION SPECIALIST MAJHAIL: All right. Let's start today's session. And again I'm Radhika Majhail with the Department of Toxic Substances Control, and welcome you all again for the second day of the GRSP. The bathrooms are still at the same location. Nothing changed overnight.

(Laughter.)

PUBLIC PARTICIPATION SPECIALIST MAJHAIL: They're still out the door, past the Byron Sher on the left-hand side. The fire exits are still the same. And just in case, of emergency, if something happens we will be meeting downstairs by Cesar Chavez Park, so -- and just a quick announcement for the members that are leaving right at lunch or right after lunch, there's a slight change in the cab program. So I hope -- the pick up location of the cabs. I hope you have the new directions. If not come see me, and then well, you know, set you up.

And without any further delays, let's start.

CO-CHAIRPERSON FONG: Good morning, everyone. A really exciting agenda for today. There are three items

that we would like to cover. The first one it's on the strategies for determining relevant factors. The second one it's the conceptual model, using the diagram the Kelly had shared with us yesterday. And actually that's going to be the if first thing we're going to cover after the public comment period.

And I think that's going to be really interesting, because Karl is actually going to go through that diaphragm, that conceptual model and give us some insight into how something like that can be used by DTSC in its decision making process.

So again, that would be the first item. The second item would be the strategies for determining relevant factors and the third item would be data gaps.

So let's start with public comments. If you have not done so, it's not too late to sign up. We do have one.

And again, I just want to remind the public that this is a working meeting for the Green Ribbon Science Panel. So the members will not be able to respond to your comments and directly answer your questions. And if your comments -- and if you're interested specifically direct it at DTSC. Again, there will be a number of workshops in which you will have the opportunity to present comments and ask DTSC questions, okay?

Thank you very much.

Kelly.

CO-CHAIRPERSON MORAN: Yes. Thank you. So we'll take public comments now, and then we might have a couple of introductory words from our DTSC leaders, before we dive into the AA related topics that Art mentioned. And we've got I think two speaker cards here. If you would wish to address the Green Ribbon Science Panel this morning, please immediately grab one of those cards from staff and put that in.

And because we only have two people, you'll be welcome to speak for at least three minutes, but I wouldn't -- I'm hoping you can keep your comments within that kind of time frame, so we keep the meeting moving. We've got a third card coming our way. And that will allow us to keep the meeting moving.

And again, I want to emphasize that the purpose of comments today is not to make comments to DTSC. It's not a general comment discussion. This is a science advisory panel, and we'd be looking for comments that would inform the Science Panel members in their discussion of alternatives assessment and specifically the items that my Co-Chair Art Fong listed for today that we'll be discussing.

So there are other forums for providing direct

input to communicate with DTSC. As they mentioned, they're having workshops. So that would be the place to do that. So with that, the three speakers today we'll start with Miriam Rotkin-Ellman followed by Will -- I think that's Lopez. Lorenz, I'm sorry.

So please go ahead. And you can move the mic to get to the right height so it's more comfortable.

MS. ROTKIN-ELLMAN: It's on. Yea. All right. Good morning. My name is Miriam Rotkin-Ellman. I'm a scientist with the Natural Resources Defense Council. Thank you all for your attention and your expertise to the topic at hand. And for the opportunity to provide some input from the public stakeholder perspective, as you all are wrestling with the technical components of the questions before you today.

And I want to speak as somebody who's relatively new to the alternative analysis field, but I think also that input is not something that might actually inform your discussion the rest of the day. And I apologize, I will be here and then be jetting out. And then I was listening yesterday via audio, because it's difficult to squeeze everything in. So that doesn't mean that I don't care, it just means that I have to go.

So just a couple of different points I want to touch on based on the reflections from the conversation I

heard yesterday afternoon. One is just from the -- in development of a robust program that the public can be come confident in, there's a fair amount that I'd love you all to consider that relates to transparency and accountability, and the development of the guidance documents. That's an essential component, in addition to the technical robustness of those documents, that will make this program a success and that will engender the kind of support from the public that I think everybody here wants.

And there were a few sort of specifics on that that came up yesterday that I just want to highlight. So there's a discussion around shifting the burden to the --what do you guys call it? -- regulated entity -- so many different words -- responsible entity. Thank you. Use the right word. And that is something that's a central tenet here and very important.

It's important to understand what is being shifted and what is not being shifted. And there's a real difference between shifting the responsibility for data, for information gathering, for disclosure to the responsible entity to shifting the decision to the responsible entity. And that makes a big difference from the stakeholder perspective.

And I just want to -- I think this Panel can

provide some very clear guidance to the agency in the development of how to be clear about the responsibilities and how they are being allocated to various people involved in this process.

Second piece on that in that sort of similar vein, a few times around people articulating different ways one could go forward. You know, people bandied around they're like we'll know it when we see it kind of idea. That, from a public, you know, raises a lot of flags from the outside world.

You know, I just want to put that reflection back in, that that doesn't provide the level of accountability and transparency the public is really looking for. I think a lot of you know that. I just think it's important to have that in front of you.

And then the last piece of reflection I just want to pull out from the conversation yesterday that I heard around was a discussion around the goal. And the goal of the program, of the alternatives analysis itself and the goal therefore of the guidance document that would then help further that goal.

And I think that's an important framework to continue to come back to. Because I heard articulated that the goal of the alternatives analysis is to identify alternatives that don't -- and then, you know, do all

1 these different things, right? I mean, that's the goal. You know, avoid risk transfer, avoid regrettable 2 substitutions. All of those are goals, in which case, 3 then your guidance document one framework for looking at 4 5 it is what incentive does that guidance document provide 6 to stop -- you know, to reach a gate where you do not need 7 to go any further in your analysis. If that guidance 8 document provides robust scientifically justified gates 9 that criteria is met by which there is good evidence that 10 the other downstream impacts are not going to happen because of those criteria, that's an incentive, and that 11 can drive a reduction in the full amount of analysis. 12 The 13 same thing around data, and data gaps.

CO-CHAIRPERSON MORAN: Thank you very much, Ms. Rotkin-Ellman.

Mr. Lopez(sic) followed by Greg Lorenz.

DEPUTY DIRECTOR WILLIAMS: Lorenz.

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CO-CHAIRPERSON MORAN: Lorenz. And I'm so sorry.

MR. LORENZ: Hello, Panel. Thank you very much for hearing me for the third time. I guess my question is about the process and some comments related to that. We certainly would like to see that the process that they use for determining compounds on the list of, let's say, whatever, it's 1100 or something, is a similar process as to what we need to provide information for evaluation.

So you talk about these matrixes(sic) of what you're going to evaluate and the critical factors, whether there's seven or whether it's 1,060 or something, you know -- and if there's data gaps we all understand that, because there are, because sometimes we're not evaluating them and you're not evaluating them. But we hope that this -- the process is such that what is inputted as to why it's on the prioritization list, is the same sort of thing is that you're looking for in the evaluation, so that we can know what the marker is we need to meet because you've said these are the things that are on the list, and here's our matrix for evaluating them.

And so it's important for us to then see how your decision making tree goes about, so that we look at future products or current products in that light. And, you know, we'd like to see that, so that it starts at the beginning in the determination of the compounds and goes all the way through the process, so at the end of the day, you know, you get our buy-in rather than you get us, you know, coming up like I did yesterday where we're a little upset.

Thank you very much.

CO-CHAIRPERSON MORAN: Thank you. Greg Gorder.

Have I got your name right?

MR. GORDER: You do.

CO-CHAIRPERSON MORAN: Thank you.

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MR. GORDER: Thanks, Kelly. Yeah. Greg Gorder, Technology Sciences Group. I wanted to comment on the discussion yesterday that -- some things I heard like Don was talking about, well, what's just concentrate on the date that are critical to the receptors that are going to be influenced. And Kelly was discussing, well, we start out with this conceptual model and so on. I mean, all that seems right.

And Helen was saying, for example, that at some point, it becomes too much, that you can't do everything. Let's do what makes sense here. And it seems to me that there's a little bit of stiffness in this guidance document that a lot of agencies interact. You know, a company could come in and say, you know, we're considering this alternative. We've done the literature review Julia talked about, for example. This is what's available. We've gone to EPA EPI Suite. We know the physical properties. Is it going to bioaccumulate? Is it going to go into air and so on, and discuss with the Agency, we don't think we have to do this, you know, and then there aren't surprises later. You know, the Agency says well, you know, we're still concerned about this or that or, no, we agree with you, and then the company has some certainty about putting the effort into going forward with that

alternative, for example. And so I just think there needs to be a little bit more flexibility than was discussed yesterday.

Thank you.

CO-CHAIRPERSON MORAN: Thank you very much.

Is there anyone else who wishes to address the Science Panel?

Seeing none. That will end the public comment period. And I think I, at this point, will turn this back over to Art.

CO-CHAIRPERSON FONG: Thank you very much, Kelly.

Let me just again go over the three agenda items that we want to cover today, starting off with the conceptual model, then strategies for determining relevant factors, and data gaps. Corey or Radhika, can you put the coal tar conceptual model slide on -- excellent. Thank you very much Radhika.

So again, we're going to have Karl Palmer from DTSC share some insight with us on how this particular model and these types of conceptual models can be used by DTSC in this decision-making process.

Karl.

BRANCH CHIEF PALMER: Than you, Art.

Yeah, if you look at this conceptual model slide, and as Kelly mentioned yesterday, this actually was ten

years in the development or it evolved over time. And it started with this framework. And without all the data there to support, you know, how significant any of these factor were, but over time they built the datasets that then inform how good this model is.

And what I wanted to do was just give you one scenario, a case study, if you will, for how DTSC used this in selection of priority products, which I think is relevant to the AA process, because our priority product profiles real are set up to look at the key factors in the regulation, just as that you have to address in the AA process.

And this was sort of not by design, but it's a good example, because one of the potential priority products you were considering was, in fact, coal tar sealants. And so the way we use this model, and if you look at this, you can see that in the upper left-hand corner of this diagram it says seal coat product, and there's a container sitting in the corner.

So when we were looking at the criteria for selecting a priority product, we did not dive in and look at all alternatives. We did not look at every chemical alternative. We did not look at, you know, every factor on the list initially. The first question was, is this product being used in California?

And so -- because part of our framework was that we, you know, if it's not being used here, or sold here, then there's no nexus with our program. So that may sound like a simple question, but there are data gaps. There's questions to be answered. And so I just wanted to highlight some of the things we did in our decision making thought process, which I think is informative.

So one of the things we first did, we said well what is coal tar sealant? What are the characteristics of it? Where are they in products? So we looked at MSDSs, we looked at products, we did some research, we talked to our counterparts at water boards, and USGS people who'd studied coal tar sealants, so we could get a picture of what that is, not at super great depth, but a snapshot.

And then we started asking the question, well, is this an issue in California? So then we looked at the market. And we have limited access to a lot of market data. And lot of the market data which we've purchased is really not designed for our focus. But we also had other tools that we used, which were, you know, do a little ad hoc market retail survey, get online see what Home Depot and other major retailers sell or don't sell, talk to contractors. So we got on the phone and we did a small survey of contractors that use sealants of this nature, and say do you have this? Can we buy this?

We also looked at the literature to say, you know, how big of a problem is this and why? And USGS has done a lot of work on coal tar sealants, and they actually have a few studies that look at where it's used, and they had a study that said that by and large these products are used east of the Rockies. And then they had another study where they were looking at some work that had been done on dust, which is, if you can see on here, is identified as one of the things in the conceptual model, and exposure pathway for PAHs.

And there was some discussion about, well, they had analyzed dust for PAHs in California and found that they were significantly lower than some of these states that had a lot of coal tar sealant use.

Now, there wasn't a definitive study. There was certainly data gaps there, but there was information that we could use to consider how relevant this was to our task, which was is this something we want to consider moving forward as a potential priority product.

So when we line all that up, you know, then there were still lots of data gaps. We didn't go through every factor. We didn't -- but we looked at the key ones, is there a significant potential adverse harm to people or the environment in California?

And our initial cut was we don't think this rises

to that level of being something we want to start with. That it doesn't appear to be a significant use. It's not to say that it isn't used sometimes. You can order it from Louisiana, you know, and use it. Why you would do that? I'm not sure.

But what we did was essentially staff did all this research. We had it on the table. People were assigned to it. And then they came to our managers and me and said this is what we've got. This is our recommendation. This is what we think. And we went back and forth on that. And ultimately in this framework, we said we don't think that fits, and we document that for -- they documented it for me. We documented it for our management. And we made a decision that, at that time, we weren't moving forward on it.

And I think that's very parallel, not at the greatest depth, but what -- the phrase I like to use in-house is, "Show your work". And I think it's very applicable to this whole process, is that there are going to be times where you get in this framework and, yes -- let's say, we did know it was used California, then we would start looking at all these different things and do the similar kind of logical process to eliminate or pull in factors that need further consideration, the presence or lack of data, comparing data if you have it or not and

how to deal with data gaps.

So this is just a mini snapshot, I think, of how we've used this type of thing. And I think it applies more generally, and we'd be interested in hearing your thoughts on that.

CO-CHAIRPERSON FONG: Karl, thank you very much. So we're going to take about an hour and have a -- an hour?

PANEL MEMBER SCHWARZMAN: Can I ask a clarifying question?

CO-CHAIRPERSON FONG: Yes. Sorry. Clarifying questions.

Meq.

PANEL MEMBER SCHWARZMAN: I was hoping maybe -Karl, thank you for that description. It's really helpful
to see how the Department used something like this. I
think I struggled with the genericness of that term,
conceptual model, to really understand how you meant it to
be used. Can you say something about how you might
picture a model like this being used in an alternatives
analysis?

So it was very clear how your staff used it in your process of investigating a product. How might it apply in the process of doing -- or of a company submitting an alternatives analysis.

BRANCH CHIEF PALMER: I think on a couple levels. One is starting with what is your framework, you know, that you're considering for your specific product and your specific business model. And one thing to recognize is that it might be different for difficult responsible entities, because whether it's the presence of your facility, you know, there are a lot of impacts that are potentially relevant for one person and not for another.

So part of it is sketching out what your framework is. And then an iterative process of how you're going to address that framework. So using the framework in the criteria in the regulations and in our -- ultimately, in our guidance that are going to help you with that, to say this is where I'm going to come back to.

And much as Kelly described that over time this specific framework led to a lot of fill-in data gaps and information that was used for probably a variety of things, that's what you would do in this process to inform the decision making supported -- or required by the AA, and then you would fill those gaps and tell your story in the AA of where it's led you, and where those gaps were, and how you dealt with them, how you considered whether they were relevant or not, how you made that determination, and I don't think there's -- I think oftentimes the perception that people have is that there's

all of these factors in the list, and that it's a checklist process. And I would say that it's not really a checklist process.

We say in the reg you have to consider all these things, but there's a default to the report being a presentation of telling that story of the logic, the rationale, the supporting documentation, the models used, and the approach. And that's going to be a dynamic thing that it is incumbent on the prepare to explain. I'm not sure if that fits --

PANEL MEMBER SCHWARZMAN: Yeah, I think that gets at it. So you're saying that this would almost be the introductory page of an alternatives analysis that would in form the Department --

BRANCH CHIEF PALMER: Yes.

PANEL MEMBER SCHWARZMAN: -- here's the way that we consider this product to be used, and what we think the potential environmental compartments are and potential exposed populations. Here's the framework for our analysis.

BRANCH CHIEF PALMER: Yes. And I think that's a good visual. And I'm a visual person, so I mean, I would love to have an AA guidance document that, you know, plop your conceptual model in here, in each box, and how it's related to each factor in the reg. And you would scroll

over, and you would click, and you would then tell your story for that. And then you could then at the end of your AA highlight your framework and show which factors were significant, where alternatives come in and how you change the potential future conceptual model, which is where we're trying to go.

PANEL MEMBER SCHWARZMAN: That's helpful. Thank you.

CO-CHAIRPERSON FONG: Thank you.

Are there any more clarifying questions for Karl on what was presented?

Cal.

PANEL MEMBER BAIER-ANDERSON: Yes. Oh, was there someone else?

CO-CHAIRPERSON FONG: Oh, I'm sorry. Julia. So I have Julia, Cal, and -- Julia, please. I'm sorry.

PANEL MEMBER QUINT: Yeah. Thank you for that, Karl. I mean it's very similar when we're like issuing an alert on something. You know, you go through the same steps. Is it used in California? And then you go through that process. But I noticed in the regulation it talks about scoping, screening, and quantitative analysis. I mean, I think those words are used at least somewhere. So to me, it's -- that's -- it's not exactly clear from the conceptual model. It doesn't translate directly to those

three steps.

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And I think that's what -- I mean, the scoping, yes, but -- and then screening would -- to me would be using something like GreenScreen or something to -- or, you know, available literature, or whatever to screen out what might be there. But the quantitative analysis that's sometimes mixed with qualitative analysis. I mean, some of that language I think we have to, if it's in the regulation, somehow this conceptual model and things that we do in the guidance document should relate to that, or explain it, or something like that.

BRANCH CHIEF PALMER: I think that's a good I think that there's some inferred -- in the point. regulation the expectation I think that for the two phases that part of -- the key part of that phase one is the workplan and scoping, so that -- to address some of the concerns that people have expressed about not getting too far down the road and going in the wrong direction, that we do view this as an interactive process, and that's a point where we would take a snapshot and say, yeah, that makes sense. Your model seems to make sense and your time frames, and you're going to have to balance out, to the extent you can at that time, the availability of data, and then the process you're going to use, whether it's qualitative or quantitative.

PANEL MEMBER QUINT: Yeah, some examples of that would help.

BRANCH CHIEF PALMER: Yes, certainly.

PANEL MEMBER QUINT: Because it's very clear from scoping that you can do the product's story. You know, what is this, how is it made, and you can get that information. But then for me even, you know, what you mean exactly about quantitative analysis and this, you know, paradigm I -- you don't know exactly what that means. So I think for some of those it would be very good to explain, you know, exactly what you mean or give an example.

BRANCH CHIEF PALMER: I hope that we can give some good examples that will inform people in the guidance. And one thing I think would be helpful for us from this Panel, there are various of you who have a lot of experience in doing some form of alternatives assessment or analysis in your business process, and in some sense, this process is just expanding the view. And when you say how do you do a quantitative assessment of something, whether it's in your realm, you're doing that for one factor or for one aspect of your business model right now.

We've added some other ones that may be you haven't done, but I think we would be informed by how

people have already done that with existing processes and apply that to these other boxes that maybe people haven't considered or that there's some ambiguity about how we're going to go about doing that.

CO-CHAIRPERSON FONG: Great. Before I let Cal answer her questions, I just got the word that Bill Carroll is, in fact, able to join us during his lunch hour. So, Bill, are you on the phone?

PANEL MEMBER CARROLL: It's total dedication, Art. I love you guys.

(Laughter.)

CO-CHAIRPERSON FONG: I know this is the kind of commitment that we want to see from our Panel members.

(Laughter.)

CO-CHAIRPERSON FONG: So, Bill, if you have any questions, please let us know, but I'm going to let Cal go first and then come back to you, if that's okay?

PANEL MEMBER CARROLL: I have none. Go ahead.

PANEL MEMBER BAIER-ANDERSON: Okay. So I think what this discussion illustrates is what Don and Kelly mentioned yesterday is that we do these -- we build these conceptual models kind of all the time automatically. And, of course, they can be narrative or they can be pictorial, or both.

And I was just wondering if it's -- if you

constructed, your staff constructed any pictorial conceptual models for the products that have been selected?

BRANCH CHIEF PALMER: No. I'm trying to think back -- you know, it was an evolution as we dove into this for the first time, but we didn't have a framework where we have a pictorial model for each one of those products. That model is essentially the narrative in the profile, which aligns with the regulation as best we could.

And you'll note that even in those profiles there are data gaps. There are things that are not there, and in some cases we called those out and said we don't know or in some cases we've said in consideration of viable alternatives. Specifically, for SPF, we highlighted that we don't know of any. But there is some research in the EU. And I think one of the commenters yesterday said there's some stuff on the horizon.

So I think it would be a good point if we did that, and that could be a starting point, and that might be something in the guidance that we could work on with the people from that industry -- each of those industries to say what is the real world model right now?

And that might be a place to start.

CO-CHAIRPERSON FONG: Good. Are there any more clarifying questions for Karl?

Seeing none.

I'm going to now open up the general panel discussion on using the conceptual -- so we're now going to open up a general discussion on the conceptual models -- wait a minute, didn't I just say that?

(Laughter.)

CO-CHAIRPERSON FONG: Wait a minute -- for determining relevant factors, and especially interested in, you know, practical experiences on the pros and cons of using the conceptual models. And I think we're going to have time-wise, yeah about 20 minutes or so. And again, in terms of making comments, please use the flag approach.

Let's see, I see Meg, Helen, Don, Cal. Okay. So let's start with Meg.

PANEL MEMBER SCHWARZMAN: My comment is just brief. I'm really intrigued with this idea of DTSC providing at the opening of the product profile a conceptual model that's pictorial, that summarizes the information that DTSC has gathered for the priority product and the pathways that the Department thinks is relevant, and then that becomes the template that anybody submitting an AA uses and makes alterations on, and adds information or says, no, this pathway isn't relevant and provides this rationale or provides additional information

about other exposure pathways or other ways that the products is used or how it's disposed of or some -- that that becomes the kind of repository of the lifecycle information about this product.

And in that way, I think it's an interesting sort of sharing of the responsibility for kind of curating that formation, that the Department puts out with the profile how you could envision this first draft of the conceptual model, and that you got a lot of feedback in a way from all of the users who are submitting AAs as sort of revisions to that conceptual model. It's intriguing to me. I think it could be really informative.

CO-CHAIRPERSON FONG: Thank you.

Helen.

PANEL MEMBER HOLDER: So I guess the question was what do practitioners do today? And so I just thought I'd answer that from what we do.

This step would be what we would -- where we'd put it in our process would be in the lifecycle thinking part. So this is what we would call lifecycle thinking.

And so just -- I said just practice would be a hazard screen first, where we look at a set of predetermine endpoints, hazard topics that we would then -- then after that would do other analyses like this. So just in terms of, like I said, practice -- the question

was what's one -- one practice is to do it that way. So we get there, but we don't necessarily start there. And the thinking behind that was to look under all the hazard rocks -- not all the rocks, but to make sure that we weren't sort of overlooking a hazard. And so we right now use GreenScreen, but you know, we could conceivably expand that, you know, for, in particular, aquatic.

But the idea would be that we do a hazard first, we do lifecycle thinking, which is I think very Similar to the conceptual model, and then based on that finding, then we could do something. And I want to be really clear, that that is the -- that the lifecycle thinking is not exposure scenarios, which is -- which happens later after this.

So that's just like I said, just answer the question of what's in practice. That's one thing. We find that that's very efficient, because we can take things off the table based on hazard, and don't have to invest anymore time, and then we can apply the conceptual model, which is relatively time efficient and resource efficient from a staff perspective to show where do we need to do the deep dive into doing an exposure assessment or doing a lifecycle.

So one thing I don't see here is the lifecycle issues that we would actually do in addition to this

conceptual model. So we would actually go beyond this in our lifecycle thinking step. Again, just to answer that question.

CO-CHAIRPERSON FONG: So I have Don, Cal, Mike, Ann, and Kelly on the list. Starting with Don.

PANEL MEMBER VERSTEEG: Okay. I want to agree with both Meg and Helen. I think this is a start. So there are multiple life stages, lifecycle stages. You know, obviously the seal coat doesn't magically appear in the bottle on the edge of the driveway for you.

(Laughter.)

PANEL MEMBER VERSTEEG: But I do think it's important that a draft be put together and be proposed put forward by DTSC, and then vetted with industry or the regulated community, because that's where you're going to learn a lot. For instance, being someone that applies this to his driveway, what this is missing is the dermal exposure, which can be significant, especially the younger the applicator. So my children come out and, you know, help me, whether I want them to or not.

(Laughter.)

PANEL MEMBER VERSTEEG: So there are other components that -- and especially when you get to the trucking of not only the container but the raw material itself in the tanker trucks before it gets to the place

where they bottle it. You know, only industry is really going to know, I would think, maybe others, what goes on at that point, and how much exposure there is and what's enclose and what's not, and how much they lose, et cetera.

So I think it's going to be an iterative process, but I think it's a good point to start with something, recognizing that any alternative you may have left something off the diagram on purpose or have a little tiny arrow going some place that turns out to be a big arrow with an alternative. So it's not -- you don't have just one that -- you have multiple diagrams for the LCA aspects, and then you have other diagrams for your alternative -- or alternative chemicals.

Thank you.

CO-CHAIRPERSON FONG: Thank you.

Cal.

PANEL MEMBER BAIER-ANDERSON: I want to respond to something Don just pointed out. I don't want to worry people by thinking that they have to generate 60 different diagrams. I think if you have a generic diagram, you can deal with similarities and differences in a narrative, too, right?

The other thing is I think, like when you see these nice pictorial conceptual models, you know, you want to -- like, well, how do we capture everything in the

pictorial?

And so on the one hand while we don't need 60 different, we may need a couple different ways of capturing different components of concerns across the lifecycle. So, like, I just didn't -- I didn't want us to go down that path thinking that everything needs a different picture. You can have one picture in variations in a text.

The other point that I want to make is just from DfE's experience, for our experience, that the -- in addition to the industry input, it is often useful to get input from the academics and the NGO community, and through that, public consultation. I think that -- that will be really valuable.

CO-CHAIRPERSON FONG: Cal, thanks very much. Mike.

PANEL MEMBER CARINGELLO: And I agree. I think it's very important that DTSC start a conceptual model. And it can be a pictorial, or as Karl was saying, that they've kind of covered that with the profiles. I think you can go either route. The pictorial ones are very helpful, and -- but I think, and I've heard this said a few times, and Cal most recently, that you need them to be iterative and have the right parties jump in and add to that.

And I want to make sure that people know it's not just DTSC who's going to put this conceptual model up. They're going to propose the initial start, here's why we think it's important. But you really have to get the responsible entity to take ownership of part of that and not just because you've got the no news, and oh, I agree with DTSC, but you've got misuses of a product that might be extremely relevant factors.

A company I used to work for, we had a product that would you never think of someone using as a consumable, because it wasn't. It was a disposal. It was an article. But as soon as we would change a formula very slightly, we would get consumer complaints. Why does it taste different?

(Laughter.)

PANEL MEMBER CARINGELLO: You know, so would DTSC have any clue that that type of product there was an oral pathway? But obviously there is, and it might be that it become a relevant factor for you to consider.

And so you need, not only the responsible entity, but I think we need to be very clear in our guidance and going out that there are other parties of interest. You know, I think we need to reach to the manufacturers of the chemicals of concern that are now linked, so the methylene chloride manufacturers, and say what -- if you look at

this conceptual model, where do you see methylene chloride, or any of these other things, where do you see them impacting that we might not have caught on to?

That you might see a pathway that is totally unexpected, you might want to look at the hazardous waste facilities and say, okay, we're looking end of lifecycle now. If you see sealant coming in -- sealant cans -- how does that interact in your facility? And we need to be able to capture that.

So I think we open the conceptual model up and maybe it's at the workshop level, and maybe the workshops turn into two days instead of one day, and say how do we make this conceptual model more robust, so that we capture all the pathways and we truly develop appropriate relevant factors. And then we go back and take our guidance and make it stronger, so that the responsible entities can put together consistent AAs, so that when it comes time for DTSC to evaluate them, they're looking at everyone starting on the same page.

CO-CHAIRPERSON FONG: Thank you, Mike. I have Ann, Kelly, and Ken Geiser next.

Ann.

PANEL MEMBER BLAKE: Thank you. So I'm trying to think of how best to display this. Maybe I'll just talk you through it, because you can sort of see that I've got

a table here. This is responding to both Kelly and Helen's question about how do we use this in practice?

And I was thinking about the most developed model that I'm currently working with. And I can share this with all of you later, but this is still very much in development. This is the framing for the BizNGO plastics score card, which we're renaming at as we speak. So it may be the chemical footprinting of plastic.

But I just wanted to give you an idea of the kinds -- of the scale of which -- pardon me -- the scale of which we're considering lifecycle impacts and how you might incorporate principles in. And I this is another way also to visualize a conceptual model of a lifecycle.

So just so you have an idea, let's see if I can remember this. So on this side --

(Thereupon a phone rang.)

(Laughter.)

PANEL MEMBER BLAKE: What key is that in Helen? (Laughter.)

PANEL MEMBER BLAKE: So this is observing a lot of work that's sort of conceptual -- capturing a whole lot of work that was done. So there was a lot of work developing the principles for sustainable plastics, and that's what's on the left here. And then across the top is the series of the lifecycle pieces of this.

So we've got polymer manufacturing, if I remember correctly -- anyway, adding additives. But you get the idea that you could take the larger bins of lifecycle that we were talking about yesterday, manufacturing, use, and disposal, and then make that a slightly more finer -- a finer grain look as you look at a different priority product.

And then I think these principles on the side are either your giant -- the big bins of impacts that you're looking at, or, you know, what is it that you want an AA to generate for you.

The reason that we had this side was to focus on, you know, we're developing the method for screening the hazard of the process chemistry of plastics, but that's a different goal. So I just wanted to give you another kind of framing of how we would go about putting a conceptual model together.

CO-CHAIRPERSON FONG: Thank you, Ann. Kelly, next.

CO-CHAIRPERSON MORAN: I'm wondering if the staff could bring up the flowchart figure. It's actually the final page of that flowchart. I don't know if you have them all here or not. This is the one -- okay. We only have one of them. Okay. Well, let's just use the one.

DEPUTY DIRECTOR WILLIAMS: Go up one.

CO-CHAIRPERSON MORAN: Go up one, please. Thank you, Corey.

DEPUTY DIRECTOR WILLIAMS: But the GRSP members should have all three figures.

matter which one. I just wanted to put that on the screen briefly to kind of point out that not -- my experience in doing these is mostly doing this kind of thing, where you wind up breaking it out into pieces and then kind of charting things through. So it's -- because I'm not an advertise, and don't think like an artist. It's actually easily to go through the different parts of the lifecycle and start thinking about that.

And of a lot of conceptual modelers will use —
there's only one place here where there's a dotted line,
but it's very common for folks to put little dotted lines.
We're not sure if something is big or small. And I
also — but mostly I want to just put that out there to
say it doesn't have to be that hard. And what's really
cool about something like this is that you can start
rearranging it and having a conversation around it. And I
just can't tell you how much I support the things I've
heard here about the idea of putting something on paper
early, and using it to facilitate conversation, because
that's the thing that — that's actually the real reason

that I was putting forth the idea of a conceptual model to help with this process.

What -- I kind of hate putting the burden on DTSC. I really like the idea of the manufacturers understanding their products, but I'm seeing an advantage to what the other panelists have been saying about getting something out there early during that regulation development process, because I think it will help people start to understand what the picture is about products and alternatives.

And I've worked with some pretty unsophisticated business types, who are really wonderful people, but aren't environmental managers, are not environmental professionals of the kind of folks who are at this table, because small companies can't afford to have the kinds of experts that are here on their staff.

And so putting something out there really helps people better understand why it is this crazy thing is happening with their product.

So the other thing I think is really important here is that conceptual modeling was new to me. Only five years ago, I was scared of the term, as I told you all the other day. So I really want to emphasize that I think it's going to be an important part of developing the practice of doing AAs, so it's a professional development

thing. And everybody is doing in their brain, at some level. I've heard I think every single person in this room talk about something that is clearly a conceptual model.

Environmental folks, people -- environmental risk assessors tend to do it more. I think Done and I and Becky and I know there's some people here who've really had a lot of experience with that kind of thing. So it's something -- this is an example of something that I think that, as a group, we're going to have to work on as we grow the profession of doing AAs. There are a number of other things that need to be worked on, but that's one.

And finally, I -- okay. Just two more minor points. It definitely changes with the alternatives. A couple people have said that. And so we need to be really careful about putting something out in the conversation for the product and not recognizing that when we start looking at alternatives, there may be -- depending on what those alternatives are, it's not just a sub -- a chemical substitute, but some other approach. It could be a very different conceptual model. And if it's a chemical substitute, there still could be some new pathways that are introduced by the chemical properties.

And then finally, the key -- my experience with this, in addition to facilitating the conversation and

identifying gaps in which it just completely excels as a tool. You can so easily get that clear scientific conversation going, which is really scientifically helpful.

It helps us prioritize where there aren't data available or where we haven't looked very hard for data, where to really press the boundaries to fill data gaps, and try to really pull the thread a little more in some area. So in looking at the whole list of relevant factors, it's too big. And so we're all trying to figure out how to narrow it down, but sometimes we need to say these are the areas that look really important and let's pull that thread a little more. Let's think about it a little harder. And that's actually what the AA process here is designed to do.

And when it comes to decisions, the same thing, a conceptual model will help us figure out which of the relevant factors probably deserve the highest weighting in the decision making process. And I know we're not at that point in the conversation right now, but the same tool plays all the way through the conversation.

So thank you.

CO-CHAIRPERSON FONG: Thank you very much, Kelly.

I have Ken Geiser, Meredith, and Becky. And then

I'm going to cut off the conversation and check in with

Bill to see if he would like to make a comment.

PANEL MEMBER GEISER: So I'm perplexed. Thanks. I'm not on.

(Laughter.)

PANEL MEMBER GEISER: I'm perplexed by the discussion. And it's interesting to me trying to sort through what I'm thinking of are the values of this, and potential pitfalls of this.

So I have to start off by sort of saying that I -- conceptual diagrams and all, I feel very comfortable with, I like. And I like graphics and all that. You know, I like to be able to see things, so I like that part of this. And when I think back to the early training we did, or the training we still do, on toxicity reduction planning, for instance, one of the first steps is to develop a conceptual diagram of the production process, so you can see the inflows and outflows and what's going on and what -- where the heat needs to be put in. Very traditional engineering kind of diagrams.

But they become the basis for then being able to look at a substitution in the process and try to figure out what's going to be impacts and all. So to the degree that this diagram here or, this diagramming, I should say, is a prelude to doing an alternatives assessment, I think it's very good.

Then I was sort of trying to listen to Karl a little bit on his description of the seal coat one, and there was this whole diagram. To me, this diagram looked like -- I don't know where the diagram came from, so it's totally out of context for me, but it looked to me like the kind of diagram you do when you're doing an exposure assessment. You're looking at all the different pathways and -- et cetera.

But I wasn't thinking we were doing exposure assessments. So that kind of derailed me. I thought this is an exposure assessment diagram, but I'm not doing an exposure assessment here. I'm doing -- mostly I'm working on hazards. And I'm -- and if, in fact, we put this into an alternatives assessment, I think it would make some practitioners think that you actually would be doing all kinds of exposure considerations in order to do this, which I don't think we've said we are intending to do.

So then I said, okay, let's get rid of the seal coat, because it's making -- it's deluding me into something else. So then I went to Meg's comment. Well, if you just do a diagram because -- if the Department does a diagram and -- it's a way of identifying what are the most relevant factors, that seemed to be important. It was a way of, well, you could do a diagram and sort of -- well these four here are more important than those three.

We're going to try to focus on that. So if the diagram is of that value, I think that is also good.

I also think the diagram could be very valuable in -- or with a conceptual modeler or however you want to say it, can be very helpful in teaching the subject of alternatives assessment to, you know, have a workshop in which you're putting up a conceptual model of how a chemical is used and how it's -- how -- what its lifecycle is, Don suggested, where it came from, where it goes to, and all of that. For some people, that's reasonably a new way of thinking. And to see it graphically can be a very explosive way to see it. They're kind of, oh, wow, I never even thought about all that kind of, where goes the waste, and all that kind of stuff like that. So I think that's really another valuable tool.

Where I think there's a danger here is thinking that this is actually part of the alternatives assessment. I don't -- I think that if it becomes part of the alternatives assessment, we're burdening -- maybe I'm just sort of reflecting on what our conversation in the latter part of the afternoon yesterday of realizing that one of the things we need to be thoughtful about is not making these alternatives assessments overly complex. And if we're asking, what I would warn us against is making a conceptual model part of the alternatives assessment

itself.

In other words, using it to build an alternatives assessment I like, but I don't want to suggest that we actually make it a part of it, because I think the message that I'm beginning to feel about the alternatives assessment is we need to be careful to be parsimonious here, to make sure that these things don't grow in scale to become some huge burdensome thing, and that has to do with the discussion we're about to have in a little bit here about what are the relevant factors again.

So, I'm sorry, I'm not completely clear in my thinking here, but those are some of the things that are going on in my mind as I think about it.

CO-CHAIRPERSON FONG: I have Meredith, Becky, and then coming back to Helen. But before I do that, since it's almost 1:00 o'clock on the east coast, let me just check in with Bill Carroll, because his lunch break is over at 1:00.

Bill, are you still there?

PANEL MEMBER CARROLL: I am, Art. Thanks very much. And I have a short comment. It's -- I've been following some of the slides. I think there may have been some other things that weren't on the website that I don't have, so I'm kind of guessing at this.

I think, you know, in a way it kind of matters

how you think. If you think in pictures, if -- generally, I think it would help you to breadboard out the process, not unlike what was shown here. I appreciate Ken's point about the seal coat thing being -- looking like an exposure assessment. And yet, depending on the situation, that may be helpful to you in understand the way a product is used.

I don't know that I have a tremendous amount to add to it. I do think that most people, when presented with this first opportunity to do an AA, I can't imagine them doing anything other than drawing out some kind of a flow diagram as to how you expect the information to come together. And the reason for bringing that up is that those kinds of things that they sort of become part of the process or part of the record may be helpful to others who are following in their path. And that's kind of what, you know, goes through my mind in building a record of these things as we go.

That's about the only semi-cogent thought that I have, at this point. Thanks for interrupting, Art.

CO-CHAIRPERSON FONG: No, thank you very much, Bill.

I have Meredith.

DEPUTY DIRECTOR WILLIAMS: I want to echo a lot of what I've heard about the iterative nature of doing

these conceptual models or using them. As you learn more, you continue to flesh these ideas out, but I also appreciate that it was pointed out that the example we used is not adequate in terms of capturing the full lifecycle of the product.

And that made me wonder, yesterday we touched on the ideas of functional use in use patterns, and we only touched on them, and I know folks have a lot of thoughts about those. And I wondered -- and this may be too big a chunk to bite off, but what's the -- so maybe it should go in the parking lot, but what's the relationship between this conceptual model and then thinking about functional use or thinking about use patterns? So it was really that question.

CO-CHAIRPERSON FONG: Becky.

PANEL MEMBER SUTTON: This is a little extension to Kelly and Ken's comment about sort of the education and training that might need to happen if we use a lot of conceptual models, which is that if DTSC were to provide an example model for a particular chemical, it might be relatively easy for folks to tweak that model for a drop-in replacement. But for something that would require reformulation or an entirely different -- an alternative that's an entirely different process or product, that gets a little more complicated. So that's just an additional

thing to think about.

CO-CHAIRPERSON FONG: Thank you, Becky.

Are there any more comments or questions before we go to the second round?

Okay. In that case, I have Helen and Kelly. And I'm going to ask you to make your comments pretty -- fairly short and succinct.

PANEL MEMBER HOLDER: So I just wanted to weigh in, in support of the idea of the Department providing that initial scoping or model or whatever it is with some safeguards though. So I think that you do need to do the refinement in workshops. That just, to me, seems like you have to do that.

The other -- there are two other things. So in safeguards, this might be something. So doing this -- doing a conceptual model doesn't tell you necessarily what hazard topics will get selected. And so there might be some value in having a baseline set that would have to be considered in all assessments.

And just some safeguards that something important doesn't get overlooked. That's Just something to think about.

A different safeguard from a practical perspective for making a compliant report. 69505.7 specifically requires that every factor, exposure pathway,

and lifecycle segment, if it's determined to not be relevant, you have to explain the rationale and identify the findings over the supporting information for that determination.

So what I would suggest that we -- that you consider is to allow, if the Department creates that conceptual model, that does an exclusion of some type, that that be allowed as a justification or a substantiation within the report. So if the Department says that we don't believe aquatic toxicity is relevant -- they wouldn't do that, but let's just say for the sake of argument that that was what they said, or sensitization, that as the assessor, when I make my report up, I can say the Department has said that this is not relevant and I don't have to redo that assessment.

CO-CHAIRPERSON FONG: Thank you.

Kelly.

CO-CHAIRPERSON MORAN: Thank you. I just wanted to make a couple of quick comments. And one is to Ken's point about whether doing these is burdensome. My experience with these is they actually simplify stuff a lot for the reviewer. And so part -- and that they're not that hard to do.

So, in fact, not only do they help you get it right as the person working on it, but for the reviewer

they're a miracle in terms of time savings. When EPA's Office of Pesticide Programs started doing these little flowcharts, in their risk assessments and their problem formulations, it's a little different thing, but environmental risk assessment, we can learn from that. As soon as they started doing these drawings, it cut review time down for me. I could scan through and immediately see if they caught the relevant pathways, if they're were thinking about the relevant factors, in terms of the things that I was looking for.

So we have to think about DTSC's time here as reviewers. That's something that's going to really, I think, save them a lot of time. So it doesn't mean it has to be fancy though. The low tech stuff works great.

And then the other to Meredith's question about functional use and use pattern. My experience is that functional use is very unique. I mean, there's a whole lot there. But use patterns, you can create generic starting, conceptual models. And there are certain use patterns where you're going to have certain kinds of exposures, so certain kinds of endpoints are going to pop up for some use patters that -- the use pattern of it's in a box, it's electronics, it's used in doors is really different than the use pattern it's painted on the walls of a building outside.

And I think eventually as this practice develops, that they'll there will probably wind up being a lot of use patterns or places in the lifecycle where there's the same thing. There's a mining conceptual model. There's various kinds of extraction conceptual models. There's things all the way through. I think we're going to start developing those things.

And then it will be easier to take the environmental fate properties, and the chemical properties and, say, for this chemical, at this point, with this conceptual model, here are the things I'm most worried about, and pull those together and roll them up in the selection of relevant factors.

Thank you.

CO-CHAIRPERSON FONG: Thank you, Kelly.

Cal.

PANEL MEMBER BAIER-ANDERSON: I just had to like react to the idea that the conceptual model can help take thinks off the table. Like I think in theory, yes, but then I worry that if sensitization isn't an issue for the chemical of concern, it may be for the substitute. And so, you know, I think strategically a long -- I agree conceptually that along the lines of if you're substituting a petroleum based chemical for -- with another petroleum based chemical, then maybe certain

things might be off the table, but you have to examine the alternatives to see if anything goes back on the table.

It's hard to -- yeah, we're all looking for shortcuts to streamline, but -- like the whole purpose is that avoiding unintended consequences.

PANEL MEMBER HOLDER: Can I respond to that?

CO-CHAIRPERSON FONG: Let's have Julia.

PANEL MEMBER QUINT: Yeah, it's related. Yeah, I was a little bit -- I'm getting a little bit confused about what DTSC's role would be initially. I mean, I agree with Meg that if you have a conceptual model, and with Ken, and the relevant factors are -- you know, DTSC says something about what they think is relative -- the relative factors are for that particular priority product.

But for the alternatives, I mean, does that mean that DTSC has to envision this with the alternatives as well? Because they will -- you know, it might be different. So I don't want to -- it's important for DTSC to maybe comment on relevant factors for the priority product. And if you have a drop-in substitute, that's petroleum based and very much like that, then the comparative, you know, analysis will be very similar.

But, you know, all holes are not barred when you go to the alternatives. And we want to encourage people to be somewhat creative or whatever about the

alternatives. So I think there's a danger of locking into that.

And I also want to comment on something that Helen said, which I think was very important in the way that their practice is or some practice that she was mentioning, of doing the health and environmental hazards first, and then seeing what drops out from that, and then proceeding, because I think we do have to -- I'm -- we're caught up in relevant factors, and I'm still trying to wrap my brain around the beginning of the, you know, health and environmental hazards, which I think are why we're doing this initially. And not that the relevant factors aren't important.

They're in the regulation. We have to do them, but I don't -- I would like to see some -- you know, how we're starting off, and then coming in with the relevant factors, as opposed to -- and nobody is saying we're leading with them, but a lot of emphasis now is on -- I guess to just make sure somewhere in the guidance that it is clear what the baseline health and environmental hazard assessment looks like, and then how this fits in as a part of that as opposed to having people go down, you know, this windy road of lifecycle with something that has some inherent hazard in it, that, you know, we want to get rid of. And maybe that's clear, but in the discussion it

became a little muddled for me.

CO-CHAIRPERSON FONG: And, Helen, you had a comment.

PANEL MEMBER HOLDER: Just responding to what Cal was saying. So could we just basically have -- start with their conceptual model and then ask what's changed?

Because that's what we do in life cycle thinking. That's exactly what we do is that we start out with, well, this is what we've mapped out for the original. Based on what we know about the alternatives, is there anything that might be different?

And sensitization is a particularly interesting one, because we have seen that as a case, right? So we got rid of a carcinogen which is fantastic. Yea. And it's a sensitization issue, and it's something that you could touch. So we actually -- that triggered -- but you had -- we didn't go back and build a new model. We took the original model and just at each sort of major thing, we said, okay, is there something different in resource uses. Is there something different here. And that -- the AB comparison, because -- and that's -- that actually, I think, works with the regs, because the regs require -- part of that four part criteria is it has to be a meaningful difference or a significant difference. And that's been our -- how we did it in the pilot and it's how

we do it in our own practice as well, is that we never go back to the ground for the alternative. We start as an AB comparison.

CO-CHAIRPERSON FONG: Helen, thank you.

I have Mike. Let's let Mike have the final comment on this particular discussion on conceptual models.

Mike.

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PANEL MEMBER CARINGELLO: And I think we don't want to lose sight of the fact that with relevant factors that when the Agency set up the profile, they gave us here is a product, here is a chemical, and here is why we picked it. So we don't want an alternative assessment to come back and say, okay, we picked this alternative, and it's better because, you know, we didn't list sensitization, but there's less sensitization with this new material. But the health impact that we picked at the -- at the start it's carcinogen. Oh, they're equivalent here, and so that carcinogenic factor is why they picked that combination. And so we found an alternative that's less hazardous for other reasons, but that first reason was the real relevant factor that caused the selection.

I think it needs to be the primary relevant factor. You can't go pick an alternative that ignores

that that was our concern in the first place, even though, yeah, I absolutely agree that you've got these other factors and it's a better player, but we haven't addressed the initial problem.

So, you know, I think there's a lot of things. I think there's a lot of good discussion here, but I want to make sure that we stick to the fact that here was a relevant factor that was addressed in the profiling, and that really needs to be something that is keyed in. And it doesn't mean that the other pieces, you know, we ignore, because I agree with what Cal said, that you don't want to pick something that there's a whole new hazard that is added in, because that didn't exist before, but we can't allow something to flow through that doesn't address that profiled relevant factor.

CO-CHAIRPERSON FONG: All right. Thank you,
Mike. That was just an excellent discussion on conceptual
factors. And I hope that this discussion was useful for
DTSC.

So I think on the agenda we're due for a break at this point, a 15-minute break. And then we'll come back and continue our discussion on alternatives analysis, following up on the second and third agenda item, which is strategies for determining relevant factors and data gaps. Oh, reminder of the Bagley-Keene requirements.

1 (Off record: 10:10 AM)

(Thereupon a recess was taken.)

(On record: 10:33 AM)

CO-CHAIRPERSON FONG: Welcome back. We're going to continue our discussion on alternatives analysis. There's again some of the things we want to cover in the remaining time that we have are, one, the strategies for determining relevant factors, and, data gaps. But actually before we do that, I understand that Lynn Goldman would like to provide us with some information about the phase one of the AA and how that connects with some of the comments that were made during the previous discussion, in terms of conceptual models and other requirements.

Yes, Meredith.

DEPUTY DIRECTOR WILLIAMS: So I just -- I want to introduce Lynn a little bit. Lynn has been the attorney who's helped us through the priority product selection, and she's been a very integral member of the team, and came up to speed incredibly fast. She jumped into the team pretty late in the game. And it's just a public opportunity to thank you for everything you've contributed, and thank you.

STAFF COUNSEL GOLDMAN: Thanks Meredith and thanks, Art, for giving me this opportunity.

Just something, you know, I've been hearing is

that you guys are saying there needs to be an opportunity to check in with DTSC about relevant factors and scoping and making sure that we're working with responsible entities to make sure that they are looking at the right things, so we don't have this gotcha situation at the end.

So I though it would be helpful to go through just the first stage of the alternatives analysis, what leads up to picking these relevant factors, and then how we're going to be looking at that.

So, you know, the first step of this first stage is that they're looking at their product, the product requirements, what they need, the functional and the legal requirements that are applicable, and do they need a chemical to meet those requirements in their product, is it feasible to take the chemical out at all or is there always going to be a chemical that are meeting those requirements. So that's kind of the first step that you have to be looking at.

And then the second step is to look at alternatives that you might know are out there, that could meet these requirements for your product. And so, you know, at that point you're already looking at some relevant factors, because you know that you need a chemical or maybe you don't. You know that there are these alternatives out there that might have other issues,

that have already been identified, because these alternatives are in the market. So you get some relevant factors there, but then you go to the third step and you're looking at whether or not, you know, factor is making a material contribution to adverse impacts.

And maybe if you're not going to use a chemical, you're really almost going to be able to skip some steps in there to say, well, there's not going to be exposure, because we can take a chemical out. So, at that point, you are already winnowing away some of the relevant factors, but if you do need to use a chemical, then you know you've added those relevant factors back in.

You're saying, okay, well, there's going to be maybe some public health impacts or there's going to be some environmental impacts.

So I think we just wanted to, you know, identify that this is only the first stage. You're identifying these relevant factors when you're thinking about alternatives, and that you're going to submit a first stage alternative analysis report, and we're going to say, well, are these the relevant factors that we were imagining, does it get to what we identified in the profile, does it get to what we know about your potential alternatives. Then we would approve that.

You go to the second stage and maybe you've done

more work on alternatives at that point, and you've identified more relevant factors. And that's why, in the second stage, you do have again identification of relevant factors. So it happens twice. And I think that's important to note that, you know, you have the first stage. But as you go through it, you get to look at it again, in case you see more of these -- the lifecycle -- sorry -- the lifecycle segments, and, you know, through worshopping, through public comments a lot of things will come out, and also through our own input on your first stage AA.

So, you know, the relevant factors does come up twice, so I think when we're saying, oh, it's hard to identify relevant factors, and maybe more will come up, that's why we are looking at it twice, and that's why there's two stages to the AA.

Thanks.

CO-CHAIRPERSON FONG: Lynn, thank you very much. And that really sets the stage for our continuation of our discussion, which is on the strategies for determining relevant factors.

Radhika, would you -- or, Corey, would you mind putting that slide onto -- again, these strategies were the strategies that we had talked about yesterday and that were offered by various Panel members. And DTSC actually

was working overnight actually got them together, and so Meredith is going to walk us through some of the concepts that we thought were -- may be useful.

DEPUTY DIRECTOR WILLIAMS: So all we did here was try to capture the conversation yesterday, follow up on Ken's quick summary -- and I'm all right really. It's worse -- it looks worse, because I tend to do that with the mic.

So I'm just going to walk through them very quickly. I'm sorry Julia that the phrase "fishing excursion" stuck, but it's a lovely shorthand.

PANEL MEMBER QUINT: Very scientific.

(Laughter.)

DEPUTY DIRECTOR WILLIAMS: But just starting with the chemical, gathering all the associated information, and working from there. You know, we could put the entire burden on the firm. Let the firm decide for itself what the relevant factors are. I used Karl's phrase of show your work to capture what that entails.

DTSC could set a minimum level, and then let the responsible entity decide whether or not there are additional relevant factors that should be considered.

Again, show your work.

I'm going to comment on this, because I think that Bob made a note on this particular one, which is in

the eyes of DTSC, of course, the minimum level is what's outlined in the regulations. And so that's -- I know that's a default and it doesn't -- it may not be satisfying to a lot of people, but, in fact, our responsibility is to use the regulations as the minimum level.

models that we discussed. We hadn't had time to dig into that yesterday. That's why spent some time on that this morning. The fifth is similar to that third one, but, you know, we do a lot of the legwork up front to identify what we think are the relevant factors. Of course, part of the challenge, as you'll see in the profiles, is there's so much that we don't know that might make it hard for us to capture relevant factors, but we could take a first pass at it, when we do the designation.

And then, of course, rely on best professional judgment and use rubrics, decision-making trees, rules of thumb to guide which the relevant -- which relevant factors need to be considered. So given that, we're hoping that now there could be a discussion of the pros and the cons of these different approaches, the strengths and weaknesses, recommendations you have or caveats about using anyone of these approaches, and what advice would you give to the Department as we proceed in developing the

guidance around this.

CO-CHAIRPERSON FONG: Excellent. And I see the flags going up already. And so what I want to do is just remind us that the meeting is going to end at noon today. And so we have perhaps -- we have until 11:30, because we're going to also spend a little bit of time talking about the next set of meetings and other types -- perhaps subcommittee work.

So, let's see, I have Helen, Ken Geiser and Ann Blake. So let's start with Helen.

PANEL MEMBER HOLDER: I have a clarifying question both for Meredith and perhaps for Lynn as well. So the original text of the legislation does have the A through M, and so you have to consider that. The regs expand that out to this extremely large list. Is it necessary that the extended list, as a minimum, be evaluated by the responsible entity, or can the Department actually assume some of that responsibility for determining relevancy.

So that A through M would be considered, but that it's -- that Ken, is that a possibility within the way the regs work? I guess that's why I'm saying it's just a clarifying question. Is it a possibility within the way the regs work for the Department to, say, decide that something is less relevant than something else and

then -- in like the conceptual model, and then have that carry forward into what a responsible entity would do, and therefore have that factor have been considered -- it was considered -- and the Department says we think it's not relevant in this case, and then have that carry through, or do the way the regs work require that every AA consider the complete set every time?

STAFF COUNSEL GOLDMAN: I think the regs do require that you look at all the factors, and I know on our part, there's a real hesitancy to eliminating any factors, because while we do know the product, the chemical of concern, we can't anticipate what the alternatives are. So, you know -- and this has been discussed that it would be very problematic for us to eliminate a factor for consideration or saying that it's been considered, when we can't anticipate what the possible alternatives are.

So, you know, I do think it would be counter to the regulations to take any of those factors off the table from any stage of the AA really.

DEPUTY DIRECTOR WILLIAMS: And then the other thing I return to is the fact again there's so much that we don't know. That we would be concerned about making statements about what's relevant and what's relevant given our knowledge gaps.

PANEL MEMBER HOLDER: So, I mean, just to sort of circle back on this great discussion we had about conceptual models though, does that take that off the table as a possibility?

DEPUTY DIRECTOR WILLIAMS: You know, what comes to my mind is what does consider mean, right? You have to consider these things. And if consider is -- I think there was a proposal that was put out this morning that said considering is developing a high quality conceptual model, and saying based on this conceptual model, you know, the following relevant factors do not appear to be significant, material. I don't know.

BRANCH CHIEF PALMER: I would just add putting on sort of regulatory hat, because what I've heard out in the community is a lot about how am I in compliance?

And I think it's -- I think, Helen, you used the term looking under every rock. What I would say is that you can look under a boulder and you don't have to look under -- and if you decide that all of the rocks and boulders in that field are not applicable, you just tell us why.

PANEL MEMBER HOLDER: That's not the question though. The question is where does the -- where can the responsibility lie? So I like the idea of the Department doing a conceptual model, but does that mean that, as a

practitioner trying to make a compliant report, that you can accept that model as being apart of your justification of why something has been excluded for consideration or do you have the obligation under the regs to go through point by point and explain in narrative form or with data as opposed to saying why something is excluded or can you just say well the Department said that's not relevant.

BRANCH CHIEF PALMER: No, I think it's the responsible entity's responsibility to do that.

PANEL MEMBER HOLDER: Okay. That changes this discussion, I guess.

BRANCH CHIEF PALMER: But I think that it's -- it may not be as heavy a lift as -- in many cases as people might think.

DEPUTY DIRECTOR WILLIAMS: So, for instance, you jumped from that to the go point-by-point narrative, right? And that -- there's some gray -- there's a gradient between identifying a category of relevant factors that based on a conceptual model may or may not be relevant to developing an exhaustive narrative for each of those relevant factors. And I think there could be categories, there could be groupings, there could be boulders that capture these things.

PANEL MEMBER HOLDER: But the responsibility still lies with the responsible entity to account for

looking --

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DEPUTY DIRECTOR WILLIAMS: Yes.

PANEL MEMBER HOLDER: -- at whatever level of abstraction that they think they could get away with?

BRANCH CHIEF PALMER: Yes.

PANEL MEMBER HOLDER: But it's completely up to them and the Department can't assume any of that consideration.

BRANCH CHIEF PALMER: Well, I think that's correct, but as Lynn pointed out, is that the structure of the regs was to have an early point of phase one, for us to affirm your assessment, and the workplan moving forward for the AA. And that may not be as soon as you would like it certainly, but I think ultimately it's on the responsible entity.

PANEL MEMBER HOLDER: Thank you for that.

CO-CHAIRPERSON FONG: Thank you, Helen.

So I have Ken Geiser, Ann Blake, Tim Malloy, Kelly and Julia. Let's start with Ken.

PANEL MEMBER GEISER: Okay. So thank you for doing that, because I did it last night myself, and it -- we almost came out exactly the same. Although, I gave them names, cute little names. And I'm more than happy to add the names to it.

But let me just step back a minute, because it is

relevant to what Helen just asked as well, and it seems to me --

CO-CHAIRPERSON FONG: Ken, sorry for interrupting. Would mine turning on your mic and/or moving it closer.

PANEL MEMBER GEISER: How is that?

DEPUTY DIRECTOR WILLIAMS: Better.

PANEL MEMBER GEISER: If I do any closer, I'll chew it.

I think, you know, haunting us in this entire work is something called no regrettable alternative or -- what's our word?

PANEL MEMBER HOLDER: Regrettable substitution.

PANEL MEMBER GEISER: Substitution. No regrettable substitution. And the reason why I think it's haunting us is that it implies that you would look at everything, because it is just possible that there might be a regrettable substitute in the one factor you didn't think about. And as I -- I was just talking to Caroline -- even to the extent of the unknown factors that aren't even things that we currently think affect us could still lead to some kind of regrettable substitute.

So obviously, there's an incentive in our work for total comprehensiveness and for total -- a huge workload. So it seems to me this question about how do

you slice out something called relevant factors is a way to back out of that in a way that is defensible, that we can kind of say we didn't look at every last thing 80 times whatever Helen's 12 lifecycle points were or whatever, but we looked at the relevant factors. And we need to find a way to tell DTSC and the responsible parties that they have a basis for determining what the responsible -- what the relevant factors are.

So I think what we have here is a set of strategies for doing that. And I like the idea of laying them out and beginning to debate them. I think this is one of the things that was making yesterday a little difficult I think is we didn't have something to look at and go like, oh, that -- well, this is more than that.

And, as I say, I'm not going to go through these, because you did such a great job of it. And I really applaud that.

I did add one more and it's a modification of this, because I'm not sure that all of these are necessarily distinct, that they can overlap, or be used differently. But the one that I think that we ought to include here is something which I might call a staged approach, in which you have a certain number of relevant factors that you look at, and then -- and I think Helen said she would look at the health impact factors first,

and then there would be another set that follow that, such that there are -- there's some primacy to which factors you look at first, and then which factors you look at second and third, because you may get a lot of diminishing returns.

If you're trying to make a decision about whether an alternative should continue to be looked at, and you can knockout that alternative pretty early by having just taking one set of relevant factors, then you don't have to do analysis on all the others. And I think that's what Lynn was suggesting a bit with that first stage, second stage idea. But it might be one more to add to this set of strategies.

But just going down through them, there's the fishing trip, there's the kind of open source, there's one in which is a DTSC base level, there's one that's kind of tailored to the products, and then there's kind of this nice thing, which you call just best professional judgment. But I like that. Please continue with that. It looks good.

CO-CHAIRPERSON FONG: Thank you, Ken.

Ann.

PANEL MEMBER BLAKE: Clearly, we have to do some education on mic technique. The singers among us can help with that a little later maybe.

So thank you. I do agree -- thank you, Ken, for laying these out yesterday and for DTSC for putting these up there, because it's much easier to have a discussion. When I immediately looked at this and started to immediately clump them and organize them differently.

So expanding on what Ken says, I think these are not mutually exclusive. I see them in two large chunks, which is kind of dividing up the responsibility a little bit. Three through five is DTSC set some kind of guidance. And I think those are different ways that DTSC can provide that guidance, different ways of visualizing them.

I say two and six, so each firm decides for itself, and/or professional judgment is going to be a little bit of something that you can work with with the industry to develop, you know, what that might look like. I think that's going to have to be a product specific piece. And number one is frankly practically how it's going to begin, how the process is going to begin.

I did want to react a little bit to something Ken just said. I know this is a discussion we've had going on about how alternatives assessment happens in practice.

Just a caution from our experience with the multi-criteria decision analysis framework is that the doing and screening with sequential factors, you can -- the hazard

is that you -- sorry, wrong unfortunate choice of words.

The concern is that you could drop out an alternative too son. So if you draw that hazard to tight. Although, I absolutely respect and I see how HP's approach has played out with the hazard screen first. So just a consideration to keep in mind.

CO-CHAIRPERSON FONG: Thank you.

Tim.

PANEL MEMBER MALLOY: Thank you.

So I've given this a lot of thought because I was really struck yesterday by Helen's pre-primal screen comments about, well, what about all the lifecycle segments and all the different factors going back? And that really kind of resonated with me, and it got me thinking about definitions.

So let me do a couple things here. One is I think it's helpful to get back to kind of the source material. And I was thankful how Lynn's comments really helped with that. But what we're trying to figure out is how to determine if something is a relevant factor or not. And the definition of relevant factor is really is it material in two different ways, right? Is it material in terms of its contribution to some harm that we're worried about? And does it make a material difference between the priority product and the alternative we're thinking about?

And it's got to have -- we've got to have both of those things, right?

So, to me, it seems like the big question is what does it mean to be material and how do you figure that out? That's what we're trying to deal with.

It also seems to me that Lynn is correct. I mean, the regulations, you know, love them or hate them, that's what's guiding everything. And they are pretty clear, in that there has to be consideration of all these factors. And they're pretty explicit that you have to explain why you dropped some of the default -- you know, the factors that appear in the regs out and why you left some in.

So, in my mind, that means there has to be consideration of everything, but that takes us to Melissa's point, which is well what does it mean to consider?

And I think that's where there's this room for flexibility. Here's my concern, right -- so let me back up for a second. So the angst that I'm feeling here, and I think it's a reasonable angst is, it comes out of two places, I think. One is that there's certain things that, you know, you'll look at certain like lifecycle segments, mining, or this or that, and it's going to be fairly obvious to you as a professional that there's not a

material contribution, right, or that it would be exceedingly difficult to collect quantitative data and do a -- you know, a rigorous scientific assessment, but you don't think it's necessary because based on best professional judgment and experience, you're pretty sure you know how that's going to come out.

And the question is -- that's get us back, so what does it mean to consider? Do you have to do the full blown, you know, turn under every rock, collect every piece of data, you know, for each of these things? And I think the answer to that has to be, no, you don't have to do that, because it won't work. I think Helen is right. It won't work if you have to do that.

So for me the big question is really, deciding what do you have to do for each of these things. I don't think these strategies answer that question? I love the conceptual model stuff. I'm like -- like I'm a visual thinker too. I love the conceptual model stuff. I don't think it answers the question.

I think -- yesterday, I said the devil's in the details. So I think the conceptual model is necessary, but not sufficient, is that what you scientists say? So for framing and thinking about what you're going to decide about, I think the conceptual model is absolutely essential and necessary. But then the hard part is once

you kind of framed it all out, is this material or not, which gets you -- that to me is the kernel question. And that's not going to get you there.

So I think that's what -- so I will go and -- I'm sorry if I'm going on, but I didn't talk at all in the first one trying to buildup some credit.

(Laughter.)

PANEL MEMBER MALLOY: I'm trying to be very concise here, because I've given this a lot of thought. So I think there's a -- I think what the -- it would be helpful, and I can't like lay it all out here, because this is just like rudimentary thinking. But here's what my suggestion would be, and I'd be happy to kind of like play with this outside, and maybe this would feed into subcommittee suggestions.

But I think it's -- there are multiple influences on this question of what material means. So one goes to this question of data availability and data cost. So the question of how far you have to go to show materiality, I think, varies depending upon the availability of data and the expense of the data or the analysis you would do with that data once you had it.

And that's what I was trying to get at. I did it inarticulately when I was just bounding on yesterday and talked about well if you're going to do that analysis

what's the big deal, you're doing the analysis anyway.

What I was trying to get at is in a world where the data is -- if there's a factor the data is fairly available and it's not -- wouldn't be that hard to do now, what I think of as your level of showing materiality would be different than a situation where there's no data availability and these regulations specifically accept the fact that you don't have to go out and develop data, right?

So, for me, that would be a less rig -- you do a less rigorous showing, but there would be more pressure on the credibility and the rigor of your best professional judgment, so show your work, that aspect of it, right.

Okay. The other thing that I think is driving it materially is also this staged thing that Lynn and -- Ken? (Laughter.)

PANEL MEMBER MALLOY: Yeah -- Ken talked about, which is it depends -- it matters -- and Ann really drove this home. It matters on how you're making the decision. So if you are doing this kind of sequential thing of knocking things out by elimination first based on certain hazard traits, that's going to -- there's going to be kind of a different level of materiality for me there, because you don't get the second -- you know, things could change later on and so on and so forth.

So this long-winded thing was really trying to

get at this notion that materiality, we should have a -- I would think that the guidance should set out kind of the -- an understanding that best professional judgment would be useful in certain circumstances where data is available. It ought to be -- which is what I think you meant, not a fishing expedition, but a sensitivity to the availability of data.

So you go out and you collect what data is available and you make judgments on those more kind of perhaps quantitatively and rigorously. Where there's less data, now you've got to fall back on professional judgment or where the cost of the analysis with data that is available might be excessive, and yet if you put ten people in a room and ask them, do you think there's going to be an issue, every single one of them would say that's not going to be an issue.

You know, so I think it's got to be this iterative type thing with a series of factors and so on and so forth.

But I do -- I'll leave with this, which is I do feel like based on what we've worried about the regrettable substitution, we can all point at examples where people thought they knew, right, what they were looking at? And it is important to make people kind of think about the factors as they go by, and then I think it

is helpful to justify why it is you went past that one, because it forces you to think about it, so you're not just doing a checklist.

I do agree with Helen that it may be that at the outset, if DTSC might be able to say based on what we know about this and the alternatives we've identified, it's unlikely that this will be a relevant factor. I think if you can give guidance in your priority listing, that would be helpful. So I think it's kind of a combination of all those.

I'm sorry for taking so long, but this is a really tough -- tougher -- like when we said here's what we're going to talk about. I thought, well, why are we going to talk about that for so long?

I guess we know now.

(Laughter.)

CO-CHAIRPERSON FONG: Thank you, Tim, for keeping your comments short. So I see that, in fact, my stern warning to you last night after dinner about -- was, in fact, very effective. Thank you.

I have Kelly, Julia, Me, and Meg.

So, Kelly.

CO-CHAIRPERSON MORAN: Thank you. And it's -I'm glad that Tim and all of us have senses of humor here,
because it's very fun to exercise them in a science

meeting, and I think keeps the appropriate tone, because I really appreciated some of the directness and honesty of the exchange that you all have and that you're not afraid of putting out what you really think. You're not -- you know, don't hold back. Tell us -- no, tell us what you think, because -- well, it's really going to help us together have that dialogue. And the fact this we don't agree is okay. And it might help us figure out a new way to go and it certainly will help the Department to hear all the different views.

So I just have a couple of thoughts.

First, is that it seems to me that some potential follow-up conversation might be merited in the areas of is this material?

And related to that, I think there's a question about is what we know important in deciding if something is material, so that this goes right to the data gap conversation that we're going to try to squeeze a little bit in and maybe take that offline too.

I'll agree with Ken, I think this whole staged approach, I'm big fan of doing that, and often do that myself. And then I really want -- I want to kind of quickly just walk through these various things. I can't -- except in the case of a fatal flaw screening -- like our company has a policy of not using any

carcinogens. So we're not even going to select alternatives that are carcinogens -- I think it would be important to do what has commonly been termed the fishing excursion.

And the important thing there is to make sure that you don't just look for a piece of data in an area, but that it's full enough that we can understand what might be the most important piece of data. And the reason I'm saying that is my experience in aquatic toxicity in particular and environmental toxicity, for human stuff there may be competing studies at the same endpoint. For environmental toxicity, there's all these different species. So we're not talking about competing studies, we're talking about actually different species that have actually different -- and there's even different endpoints, so it's not just it dies. We're also wondering about does it grow and reproduce, can it still swim, is it going to be able to -- will its eggs hatch, you know, all these other things.

So stopping -- I think some of the methodologies that are out there say, oh, find your Daphnia magna LC50 and your fathead minnow LC50 and you're done. And in my experience, if you don't look for the rainbow trout and the Ceriodaphnia and you don't start looking at all the other species that are out there, you could miss something

really important to decision making.

And maybe none of those things exist, but if -- I want to make sure I emphasize the importance of being completely enough. I'm not fond of each firm decides for itself. I think that would be extremely hard for the Department. And I've also had some experiences where -- in certain instances, where you decide for yourself, you're going to decide for yourself in a way that favors the decision that you'd like make at the end, because you might have an economic interest in one decision versus another. And maybe the customer doesn't have it, but maybe someone other than the customer is doing the AA, and that's probably going to happen.

I'm also not very fond of DTSC being asked to set some requirements, but I do think that it makes sense, and I think it's already, in fact, in the regs that there's kind of a minimum level of things that have to be thought about. So I'm not sure that's a special separate thing.

As you know, I like the conceptual model in terms of helping move that forward, and particularly helping us know what we don't know that might be important. So that's it.

And then I've got to say I'm really down on best professional judgment, and for the reason that Tim kind of got at. All of us can name time after time where you had

a group of really smart people who got together who were just convinced that there wasn't going to be harm from something, and they were wrong. And my favorite case of that is PCBs.

So I -- one of my latest things is PCBs and building caulk. So we know it wasn't going to be there. It's not going to be there for pesticides. It's pet flea treatments. We're going to put a spot of something on a pet and it just disappear from the world. And now we know that's not true either.

So those are just examples of best professional judgment by good smart professionals can still be wrong. So I think that the other more scientific approach is where we pull that thread a little bit and think it through a little more are important.

Thanks.

CO-CHAIRPERSON FONG: Thank you, Kelly.

So I have Julia, Meg, and I'll the final comment on the strategies for determining relevant factors before switching gears over to data gaps.

Julia.

PANEL MEMBER QUINT: Yeah. Julia Quint. Yeah, just -- I think, you know, what's really important is to -- for DTSC not -- to set -- to define an approach to looking. I mean the fishing excursion, you will have to

search the literature or whatever you do, I mean, and look at physical chemical properties, and try to predict, look at structure activity for the alternatives, you know, or whatever, to try to predict what is important in terms of, you know, health and ecological hazards, and -- as well as all of the lifecycle information.

But I think, you know, having some idea of what approach you should take it will be important, because I think somewhere in the regular it said, you know -- it was very vague about where -- looking for information -- best available information or available information. That could be Google for some people. It could be a variety of different things PubMed for other people.

So I think having some sort of structured way of doing the fishing would be important. You know, if you're a soil chemist, what would you look at in order to rule out soil contamination or something like that. I'm not a water person, but what would be the logical places for you to look for information for a chemical if you were trying to rule out effects on water or something like that.

There has to be -- I mean, that's happened for other regulations. Like for the Hazard Communication Standard, they would have a series of references or places for you to look for the information. If you've exhausted all of those, then at least you've made some attempt to get the

information. If you leave it very vague, you know, best professional judgment can mean anything to anybody.

So I think it would be important if -- to the extent possible to think about the approach you would take if you were trying to find out information about a chemical that you didn't have -- you know, that wasn't well studied where it's not data rich, and what would that approach be? And then that would be a way to evaluate the AAs. If the person has gone through and systematically looked at all of these different resources and there's no information available, either the alternative can't be used or at least you know that some attempt has been made to, you know, assess the hazards.

And that's all we do in government. If you're trying to look for something and do a comprehensive evaluation, you just look, you know, at all the sources for that information. And if you don't have it, then -- you know, and we're not asking people to develop it. So I think that would be important though in the guidance is the approach to looking.

CO-CHAIRPERSON FONG: Thank you, Julia.

 ${ t Meg.}$ 

PANEL MEMBER SCHWARZMAN: Thanks. I wanted to bring up briefly the document that Helen gave us last night and circulated, because I think my favorite also

from the list up here is the chunk that's sort of three through five and some variation on that, where, you know, the combination of the conceptual model that DTSC puts forth at the beginning, sort of says here's what we think is important. Now, all you stakeholders out there, either responsible entities or public health scientists or NGOs or whatever, what have we missed?

And that gets, you know, added onto and iterated on. But then that's just the starting place, and paired with -- with the, you know, type of kind of guidance that Helen handed out last night, I think it's potentially an approach that firms can use -- responsible entities can use that helps start to answer the question what is it to consider, the A through M?

Sort of does it go down the drain? Okay. If so, what effect does it have on water? And obviously, it's not the complete how-to guide, because it doesn't have -- it doesn't address Julia's question of what does it mean to answer this question what is it's affect on water? So there's still some steps about how do we -- what steps do we expect responsible entities to take to be able to answer those questions, how far do they have to look, in a sense?

But I like it, because I think it's the first -it's a great starting framework, so you take the

conceptual model and then you run through all these questions. And you need a little bit of guidance about how to -- where to look to answer those questions.

The one other piece that I wanted to bring up is something we haven't talked about yet at all, because it hasn't really been particularly relevant, but remembering that this AA goes into a regulatory decision. And remembering the range of actions that DTSC has at its -- in its toolbox to take. And so there may well be a big question mark on some piece of the conceptual model, or, you know, missing data. This is partly to help you make the transition, Art, to the next section.

That I can imagine some creative responses that DTSC could have in terms of the regulatory actions that they take in response to these submissions. And it could be some combination of actions that pairs filling data gaps with actually making some substitutions, or making a substitution that seems to make sense based on the information available. And, by the way, would you please monitor this effect that this has on water, you know, for the next year and get back to us about it? Or that there's -- that there's those kinds of -- there's such a colorful palate of actions that DTSC could take that could be formed from, you know, the questions that arise because of the conceptual model and the way that responsible

entities cruise through it. I just wanted to bring in that element.

CO-CHAIRPERSON FONG: Thank you. Before I make my comments, I just want to do a time check, because people have flights that they need to get to. And it's almost 11:15 and our meeting is scheduled to end at 11:30. And we're going to spend the last half hour planning for future in-person meetings and possibly subcommittees. So should we continue with this conversation or just --

DEPUTY DIRECTOR WILLIAMS: I don't think that we would do any kind of justice to a conversation on data gaps at this point. It's just too big a can of worms to open at this late hour. And so I think -- and this conversation is valuable, so just a few more -- you know a few more minutes on that, and then we could move to next steps.

CO-CHAIRPERSON FONG: That sounds great. So I have Helen, Ken, and myself. Helen.

PANEL MEMBER HOLDER: So I kind of have another clarifying question again for Meredith and Lynn. So looking at these options or strategies potentially, it actually -- it's kind of in response to Kelly's distaste for number two. Number two is actually the only thing that's supported by the regs, right? So I just want to

make sure that that's clear. It's like -- that these other things -- anything that involves DTSC making some narrowing, that's the question that we asked before, right? So number two is actually the only thing that the regs will support, is that true?

DEPUTY DIRECTOR WILLIAMS: In the narrowest sense, yes, right.

BRANCH CHIEF PALMER: Well, 1 and 6 would also be supported.

DEPUTY DIRECTOR WILLIAMS: That's true.

PANEL MEMBER HOLDER: Okay. Okay. So the firms -- so the responsible entity is on the hook for doing all that. That's what we established before.

DEPUTY DIRECTOR WILLIAMS: Yes.

PANEL MEMBER HOLDER: And so I guess my -- if that's true, then the question that I have is can the Department give any guidance at all on determining relevancy? Because anytime you start weighing in on what you think could be eliminated, you're endorsing an incomplete consideration of the set of factors. So is this all a moot question for guidance and for mapping that initial model?

BRANCH CHIEF PALMER: I'll add, and then Lynn you can jump in. But I think, you know, we've always had issues of where you draw the line between guidance and a

rule of general applicability or a determination. And I think what we're hoping to do is give examples, scenarios, tools, and framework discussion that will inform the responsible entity in the making of the decision in the AA.

So if you're asking can we, by -- essentially do a de facto exemption or exclusion of some factor that's required in the reg, the answer is no. Can we frame it and say this is our thinking, and this is where we suggest you look, yes. Can we -- and then when -- the other thing I want to stress is that our hope is that these are not going to be processes that are done in a vacuum, is that as we go through the process of having the workshops and dealing directly with responsible entities doing the AA, that there's an opportunity for dialogue and discussion to help people. So that once we get through guidance development, which is our focus, is that we'll help people through it.

But my angst is that ultimately, it's the responsibility of the entity. We want to do as much as we can to help that decision-making process, but we can't be the decider at that point in the process. So cutting something off, no, but we could suggest, for example, in a framework saying this is our perspective of why we chose this in the priority products selection. Here in the

guidance, if we want to do specific guidance on a specific priority product, we could say these are our questions that we think are of interest.

But a lot of it is we don't know the answer or the lack of data or not and that the framework is asking people who know more about it. So understood that's not -- there's a tension there.

STAFF COUNSEL GOLDMAN: Just something -PANEL MEMBER SCHWARZMAN: Could I just do a
clarifying thing, because I felt like what I said was
misunderstood, maybe about three through five? Is that
all right? And then I would put my flag down. That's
all.

CO-CHAIRPERSON FONG: Yes, please.

PANEL MEMBER SCHWARZMAN: Okay. I think I was picturing it the way Karl just described it, not that three through five describe a narrowing that DTSC does, more like a pointing in this direction, and then using a worksheet, for example, like your factors that step a company right through A through M. They consider all of those. It doesn't -- it doesn't -- DTSC's pointing in a certain direction doesn't remove any of the factors, A through M, from consideration. That wasn't my understanding of options three through five up there.

STAFF COUNSEL GOLDMAN: Yeah, so I just -- I want

to agree with that, that really what we're saying for relevant factors, what you need to consider. One of the things you have to consider is the exposure factors that we considered when we prioritized, so the things that we put into the profile, so kind of echoing both Karl and Meg.

We're not going to take anything off the table, but we certainly do highlight things that we think are absolutely relevant. So there is kind of a minimum data set there, but we wouldn't take anything off the table.

PANEL MEMBER HOLDER: And I think that this is a really important point for us to discuss, because there's a difference between deciding something is important and taking something off the table. There actually -- although, it results in sort of the same decision, I mean, the question that I keep having is -- in my mind, is how do I justify excluding something? What is the substantiation? What level of consideration do I have to show in order to exclude?

And I guess the reason that I was asking this question as a procedural question was, it may be -- I guess my question was can the Department, even in a guidance document, give you any indication of what can be taken off the table? Is that allowed within your authority to even give that sort of a guidance, that it's

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okay to not look at a factor under certain circumstances?
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             DEPUTY DIRECTOR WILLIAMS: And I think we then
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    run this risk of underground regulations or, you know
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   being in --
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             PANEL MEMBER HOLDER: That's exactly why I'm
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    asking.
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             DEPUTY DIRECTOR WILLIAMS: Yeah. Doing something
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    that explicit in the guidance is very likely to be counter
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    to the regulations.
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             PANEL MEMBER MALLOY: Can I just jump in for just
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    a second just on the legal question? It's like the only
    time I get to talk with any authority.
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             (Laughter.)
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             PANEL MEMBER MALLOY: And even that is moderate,
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   because I'm not DTSC, but could I just say one quick
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    thing?
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             CO-CHAIRPERSON FONG: Can I stop you?
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             (Laughter.)
             PANEL MEMBER MALLOY: You can stop me now.
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             (Laughter.)
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             PANEL MEMBER MALLOY: I just wan to say --
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             CO-CHAIRPERSON FONG: We always appreciate your
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    insights.
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             PANEL MEMBER MALLOY: Okay. And I'm not jumping
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    in here reflecting that I support the idea that DTSC
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should or shouldn't do something, like what Helen is saying, but I take her question as being more narrow than what's being answered, because I take her question as if DTSC were to make a statement say in the priority product listing and they'd identified the priority product and a few alternatives, and said we've looked at all the data and based on our review of available data, we don't think, say, there's a greenhouse gas issue with any of these or that these come from the same manufacturing thread and therefore you wouldn't have to look at that lifecycle segment.

The question is whether in complying with the regs, the responsible party could say we're dropping out this segment and we're relying on the data -- the statements the DTSC and the supporting data that they use, would it be a -- would that be a way in which there'd be this iteration, this interaction?

And, to me, that seems like -- whether you agree or disagree with DTSC being in the business of doing that, as a legal matter, I don't see that there's any legal barriers to that. It's not an underground regulation problem, because that's in the priority product listing that went through the regulatory process, right?

So I'm just saying, and its -- to me, it seems like there's legal authority to do it. Whether it's a

good idea to do it, you know, I don't -- you know, that's not my call.

CO-CHAIRPERSON FONG: Thank you, very much, Tim. I have Ken and Don.

Ken.

PANEL MEMBER ZARKER: Sure. Ken Zarker.

So a very helpful discussion. I thought there's a lot of good ideas here in terms of how you might approach this. You know, I think more of maybe a hybrid approach.

But I go back to a couple thoughts. We used to, you know, talk about the letter of the law versus the spirit of the law. And I feel like this conversation is going towards a really compliance oriented approach. And my fear with that is that people will look at this as a check box exercise. They'll meet the conditions. The staff will review that. So the missed opportunity here. And these things are also going to be snapshot in time based on available science and the data.

So I've been thinking about how DTSC might promote or what we could learn from product innovation. So how do organizations go about doing product innovation?

In the old days of pollution prevention, we got a multi-disciplinary team together to address these issues. So I think these are all helpful ways to go about a

process, but you might think about putting guidance on how you might organize a team or a process that promotes product innovation, because I think that's, at the end of the day, what we're trying to advance through the science.

CO-CHAIRPERSON FONG: Thank you, Ken.

Don.

PANEL MEMBER VERSTEEG: Yeah. I just wanted to quickly say something in favor of best professional judgment. This will get into data gaps, but sooner or later you're going to have a glaring data gap. And you can do structural alerts, a.k.a. best professional judgment, you can do QSARs, a.k.a. best professional judgment, or you can do best professional judgment.

So, at some point in time, you're going to have a data need that you're not going to have data to fill and you've got two choices. You can, you know, use best professional judgment and all the tools that are very similar to best professional judgment, or you can run a study.

I'm not sure -- I don't know if the regulations allow you to go off and do studies if there's sufficient time to do that in the regs or if that's even supposed to be part of this? But best professional judgment is going to have to be used to fill in data holes.

CO-CHAIRPERSON FONG: Thank you very much.

And I will end today's session by making a comment that's actually it's going to tie in Ken Geiser's comment on a staged approach and Ken Zarker's comment on innovation. So, you know, as I understand what you were getting at Ken, was in terms of the stages approach, you would have kind of an initial set of relevant factors that would help you rapidly screen out certain alternatives.

And the initial set may be something like, you know, the most -- the major -- important relevant factors, such as human health impacts. And I think that's a really good approach, but I think it's premature to use that by itself, because -- I'm sorry to do that by itself without taking on considerations of use, exposure, performance, availability, and other factors, because that goes back to Ken's comment about innovation.

In order for us to do innovation -- so let me give you a very specific example. Right now, the semiconductor industry is trying to figure out how to make semiconductors smaller, faster, and more energy efficient. And one of the pathways for doing that is to use certain types of compounds, such as gallium and indium. And if you were to use the staged approach without taking other factors into consideration, you would pretty early on eliminate that as a possible way of moving us forward.

And that really -- it's -- if you were to try to

do that in the semiconductor industry, it would just really be a major roadblock to the innovative process. So just something to keep in mind, or just want to make that point.

So I'm going to now switch the mic over to my co-chair Kelly Moran who is going to walk us through the next steps.

CO-CHAIRPERSON MORAN: All right. Well, I'm probably going to fairly quickly turn this over to Meredith. But we're at the point in the meeting where we need to figure out where to go from here. We've had, I think, a really amazing set of dialogue over the last day and a half. And you all have individually contributed in all kinds of various ways a lot of very stimulating thoughts that I've personally found very exciting. And it's just the start down this part of journey towards helping the Department through its journey in implementing this regulatory process.

So our goal in the next half hour is partly substance and partly logistics. So I'll be asking Meredith to start off by saying whatever. I know you'd like to be thinking about what next steps there are, and you may want some input from this Panel about topics that we might want to consider.

But I think the most important thing is what does

the Department need, because some of the things we're talking about here today have been conversations that the staff have been having for months, and aren't surprises. And so it's not just because we're sticking on something doesn't mean that the staff needs more from us in a particular area. So I want to start with Meredith, and I think she's going to ask for some advice, but our focus in the next steps should be what is the Department going to need?

So, Meredith.

DEPUTY DIRECTOR WILLIAMS: Thank you. I would -that was a great set up, simply because there is this
element in which a lot of the topics that we're discussed
have been discussed by the alternatives analysis team.
And yet, the great thinking, the clear articulation, the
varying perspectives really take us to the next level in
terms of being able to really practically develop some
guidance, and capture things, knowing what needs to be
captured in the guidance, whether that's a more adequate
discussion of what material is, or whether it's examples
of how you might use conceptual models, et cetera.

I think we walk away with a lot that's really quite actionable. And so even though some of you are frustrated or confused, the vast majority of the conversation was directly relevant to the work that we're

doing. So I really appreciate all your thoughtfulness.

In terms of next steps, one thing we had thought about doing, and I'm just going to throw out one or two topics that could be considered by the GRSP, and then hopefully there will be an opportunity for Panel members to make suggestions. But this issue of data gaps is not -- it's going to go away.

And so there are questions around how do you decide whether there really is a data gap? What's the hierarchy of data knowledge from, you know, an authoritative list all the way down to best professional judgment, and everything in between? And at what point do you say, yes, this is a data gap? And then you have to decide whether that data gap is material, and then you have to decide what you're going to do around it.

I'd be very interested in hearing, especially from the practitioners, what they have done to make those decisions? Is there a data gap? Is it material? What am I going to do about it? So that's a topic that I think we would really like to hear the Panel explore in the future.

I know Ann is probably going to say a little bit more about functional use. So I'm not going to refer to that one so much. Bob, this is your chance to -- do you have anything you want to just throw out as a concept?

SENIOR HAZARDOUS SUBSTANCES ENGINEER BOUGHTON:

Well, one of the other overarching elements that's been brought up, we haven't really talked about decision making, but trade-offs was talked about and how do we give guidance to people to consider trade-offs and decision making? So that's another similar to what is relevance, very similar.

DEPUTY DIRECTOR WILLIAMS: Yeah. And in the long term, of course, as we get closer to a regulatory response, you know, how do we cultivate transparency around our decision-making process, what tools are useful for doing that? These are the things that maybe are not on our front burner, but certainly are going to be coming at us pretty quickly.

So those are just a couple topics I'd be interested in hearing whether or not the GRSP thinks those are good topics to pursue, and I'd be interested in hearing additional topics also.

CO-CHAIRPERSON MORAN: So I see Ken Geiser and Ann Blake, one, two, and both of those questions on the table.

Ken.

PANEL MEMBER GEISER: I think the topics you've suggested look good. Those are ones -- obviously, we were going to try to get to data gaps today. We didn't, so -- and I do think that's appropriate.

The functional use thing I think is very interesting, and California could really do a good job at helping to ferret out a longer discussion that's going on across the country, so I really like that.

I do want to put a plea though into something I said yesterday, which is I would hope -- I really feel it's important that DTSC spend some time looking at how they're going to actually evaluate the alternatives assessments they get in. And I feel that to wait too long on that is going to be problems down the road. So maybe you don't want us to help you with that, but here's just another moment for me to make that plea. If the GRSP can be helpful in that, I would urge you to think about that as well.

CO-CHAIRPERSON MORAN: All right. Ann followed by Meredith and Becky.

PANEL MEMBER BLAKE: Actually, since Ken has said that I think I'd want to echo that as well. That came up a lot yesterday but definitely you can't really forward without -- with guidance without knowing what it is you're going to be evaluating for. So just to put another plug in for that.

I wanted to give a little more substance to the topic of functional use and to say that while there is an ongoing conversation -- and Cal is very much a part of

this as is Ken, so feel free to add to that -- I'm not sure that this is necessarily the time you want to think about functional use.

So where the conversation is now, in the broader practitioner community is thinking about what kinds of information do we need in a functional use, how do we categorize functional use, and what kind of information is it, that we need to make a decision -- a good decision on an alternative, so to avoid regrettable substitutions and so forth.

So that's where we're at, what's the granularity of information that's necessary, how do we find that, where are the data gaps, kind of -- it's a parallel conversation, but I think it belongs more in the alternatives assessment when you start getting evaluations of alternatives in

CO-CHAIRPERSON MORAN: So Meredith, do you want to jump in now? I see your card is up.

DEPUTY DIRECTOR WILLIAMS: Yeah. Another topic that we touched on yesterday, but we didn't -- and I know that all of you have lots of opinions about whether or not these were color coded properly. And we will just take for granted that they're not colored properly.

However, there are general patterns in this table that we looked at that we didn't really dig into very

much, which is there are certain areas where we know that there are -- there -- the frameworks that have been developed to date really have not delved deeply into those particular areas, and they are required to be considered as part of our regulations, and therefore there's a gap in the science, there's a gap in the state of the art when it comes to alternatives analysis.

And there's a question of, you know, what can DTSC or California due to encourage the community of practice to start digging into those things? There are questions -- this raises questions about, you know, what we're asking responsible entities to take on, given that we know already that in terms of water conservation there's not a lot of depth of knowledge out there or not a lot of -- the frameworks don't go as deep as we'd like them to go.

So that's another thing to think about is how do we fill in the gap in the kind of the state of the knowledge?

CO-CHAIRPERSON MORAN: Becky.

PANEL MEMBER SUTTON: I'd like to do a little bit more discussion on situations where we're not dealing with drop-in replacements, because I think that might be more the exception than the rule, if you can't just eliminate and unnecessary chemical.

that something that may or may not be appropriate here, but I'll just throw it out there, is about professional development. The last time this group was together there was actually an assessor requirement and so forth in the regs, and that fell out. And I know there are some other movements afoot on professional development. So I don't know if there's need for a conversation here, but I'll just throw that out there as something I keep seeing is how is that conversation going to happen and how DTSC relates to that, and the fact that many of those conversations are around something that's a little smaller in terms of its capacity than what we're talking about here, so -- and I just saw Cal stick her card up.

PANEL MEMBER BAIER-ANDERSON: Well, I think on that note this field is still evolving, so even when we discussed the conceptual model, the conceptual model for risk assessment may mean different things for alternatives assessment. And so, you know, having or adding to the guidance materials definitions and an explanation of how we can appropriate tools from other disciplines to help, but how that would be different from the standard use, I think, could be useful.

But again, getting back to this concept that it's evolving as -- you know, as we're meeting, this concept of

alternatives assessment is evolving. So that makes it tricky, but it also makes it critical that we have some type of development process.

CO-CHAIRPERSON MORAN: I'm not -- oh, Helen, I missed you.

PANEL MEMBER HOLDER: It was in our parking lot, but I'll put it on our next steps as well, is substantiation levels. What do you need to do to prove that something is excluded, as relevant? What do you need -- what's the documentation level that you need to do?

CO-CHAIRPERSON MORAN: And I'll take the Chair's privilege of attaching to that, Tim raised the definition of material. It's kind of a related thing.

I don't see any other flags up, at this point, so have you heard what you needed from this group right now?

DEPUTY DIRECTOR WILLIAMS: I've heard. That's quite a range of topics, and so we will have to do some thinking about priorities and -- yeah.

CO-CHAIRPERSON MORAN: Okay. So in terms of next steps, how do you want to proceed? We have about 20 minutes. And our thought had been to try to think about doing some discussion of scheduling and what kind of meetings we might want to have. Do you want to go ahead and do that at this time, or are there other things?

DEPUTY DIRECTOR WILLIAMS: Well, the short reason to have that discussion is that our current thought had been to have a another meeting in the fall. And if we have a meeting in the fall, that's after the -- after the workplan will be published, so there wouldn't be an opportunity to weigh in on the workplan. And then we'll be in the home stretch on some level of guidance. And so it's a little bit late.

So it would be valuable to the Department, either to have subcommittees that we can engage with in the intervening time frame or have at least a conference call meeting to talk about the workplan and talk about, for instance, this issue of signaling the endpoint about our decision-making process, and bringing that back before the Committee and getting some input.

CO-CHAIRPERSON MORAN: All right. So do you want to think about any of that scheduling right now, or where do you want to go. We have 20 minutes. We can do something with it.

DEPUTY DIRECTOR WILLIAMS: Well, I did -- yeah, I did see some nodding of heads that people seem receptive to the possibility of having something this summer?

(Head nods.)

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DEPUTY DIRECTOR WILLIAMS: Yeah, anybody got a -- I mean, -- okay. Great.

Corey, do you want to -- I'm going to look at you in terms of whether you want to try to tackle some scheduling now. And at the very least perhaps we should look at the October dates anyway and ask people if they're available on the October dates, which they don't have. Do they have?

We have potential dates, September 2nd, 3rd, and 4th. We're not asking you to come for three days, but those are -- within those three days we might ask for another day and a half. And I'll tell you that's the day -- those are days right after Labor Planning. So if you're planning to go away on Labor Day, that might not be ideal. October 7th and 8th, 13th through 16th, October -- PANEL MEMBER SCHWARZMAN: Can you slow down and repeat those.

DEPUTY DIRECTOR WILLIAMS: I know. I'm just -- I got warmed up. I just had my own copy.

Let me know when you're ready for me to -- okay. So I said October 7th and 8th, October 13th through the 16th, and that's the Monday through the Thursday of that week. October 20th and 21st, that's the Monday, Tuesday, and October 29th and 30th. I'm sorry. Do we get Columbus Day off?

BRANCH CHIEF PALMER: No.

DEPUTY DIRECTOR WILLIAMS: Sorry, just making

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So that said, I'd love to hear known conflicts first. If people have known conflicts with any of those dates, please let us know.

Yes, Corey, so we have some known conflicts here.

PANEL MEMBER ZARKER: The last week in October is problematic for me. We have our Green Chemistry

Roundtable meeting up there in Seattle.

DEPUTY DIRECTOR WILLIAMS: Okay.

CO-CHAIRPERSON MORAN: Maybe an easy way to handle that would be to just since I think everyone got the dates down -- did everyone get the dates down? Maybe we can just go around the room and ask if you have a conflict on any of the dates, and maybe the staff could keep track of those.

DEPUTY DIRECTOR WILLIAMS: Okay. Don, do you know of any conflicts.

PANEL MEMBER VERSTEEG: Last week in October is bad.

20 DEPUTY DIRECTOR WILLIAMS: Okay.

21 PANEL MEMBER SUTTON: I have a conflict on 22 October 14th.

DEPUTY DIRECTOR WILLIAMS: Okay.

PANEL MEMBER SCHWARZMAN: I think I'm okay,

25 | except for teaching, but that's always there.

1 PANEL MEMBER QUINT: None that I'm aware of.

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PANEL MEMBER MALLOY: I have a problem with the 7th and 8th, we have like a big conference on the 9th and the 10th.

PANEL MEMBER HOLDER: I was going to say no conflict, but I guess whatever he's got, I probably have that too.

PANEL MEMBER GEISER: Okay. So the first week of September I can't do, and the last week of October I can't do.

DEPUTY DIRECTOR WILLIAMS: Okay. Well, let me just tell you the only date I've heard so far that works for everybody is the 20th and the 21st. So don't blow it. Yeah. So, Mike, what do you got?

PANEL MEMBER CARINGELLO: The only conflict for me was the last week of October.

PANEL MEMBER BLAKE: Same as has already been -- Tim's conference on the 9th and 10th, and the last week of October.

DEPUTY DIRECTOR WILLIAMS: Okay.

PANEL MEMBER BLAKE: The last week of October is an AA Community of Practice, so that's going to take quite few of us.

DEPUTY DIRECTOR WILLIAMS: Okay. That's not going to work.

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Cal. 1 PANEL MEMBER BAIER-ANDERSON: I'm good. 2 3 DEPUTY DIRECTOR WILLIAMS: Okay. So I think we 4 actually -- oh, no, you guys don't count. 5 Just kidding. I'm sorry. 6 CO-CHAIRPERSON FONG: I'm going to take the 7 positive approach and tell you that I'm open on the 20th 8 and the 21st. 9 DEPUTY DIRECTOR WILLIAMS: Okay. 10 CO-CHAIRPERSON MORAN: Me too. 11 DEPUTY DIRECTOR WILLIAMS: So, Corey, those 12 are -- by the way those are dates when the Citizen is 13 available. So we can go ahead and reserve a block of 14 rooms and start thinking ahead to that meeting. So we'll 15 just presume that that's going to be the day. Yes. 16 PANEL MEMBER MALLOY: Two other members who 17 aren't here today --18 DEPUTY DIRECTOR WILLIAMS: Bill and Julie. 19 CO-CHAIRPERSON MORAN: Yeah, Bill and Julie. 20 DEPUTY DIRECTOR WILLIAMS: Very good point. Ιn 21 which case then, I should ask you to not just do the 22 affirmative, because if we come down to --23 CO-CHAIRPERSON FONG: In that case, October 7th 24 and 8th is our and October 29th and 30th is out.

DEPUTY DIRECTOR WILLIAMS:

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CO-CHAIRPERSON FONG: Thank you.

PANEL MEMBER SCHWARZMAN: I just wanted to point out, I don't think you needed to scratch the entire October 13th to 16th section. It was just the 14th Becky was unavailable.

DEPUTY DIRECTOR WILLIAMS: Yeah. Okay.

CO-CHAIRPERSON MORAN: So we could do 15th and 16th. I'm looking for calls on October 15th and 16th. Okay. So all the heads are nodding there. So we have two sets of possible dates to check with Julie and Bill. And so hopefully one of them will work for both of them.

So I think we're good there, but we would also be anticipating having one or more conference calls this summer. And those we'd probably schedule -- since those don't involve travel, they're a little easier to schedule. So but that -- it might be a conference call of more than a couple hours. If we're going to get this together, we might spend a half a day or more. So we want to make sure folks are going to be committed to doing that?

And if we have subcommittee calls, our general agreement is that if we have an official subcommittee, we would bring that back to the full GRSP to at least talk about the recommendations. So I just want to keep that in mind. I don't know if we're going in that direction. But if we wind up in that place, that's something that

everyone needs to kind of remember and think about, that it is going to be important to be prepared to participate in those.

DEPUTY DIRECTOR WILLIAMS: I'm going to jump back to possible topics for future discussion, which is that the National Academy report might be out by October, and that could be an explicit agenda item that we -- you know, we hear about that.

CO-CHAIRPERSON MORAN: And if after this meeting someone comes up with something that they want to suggest is the best thing to email Corey, since she's our lead contact.

PANEL MEMBER GEISER: Just on the conference call, can you avoid August?

DEPUTY DIRECTOR WILLIAMS: Yeah. I'm already kind of anxious about summer. You know, so we'll see what we can do. And actually for us earlier is a little better because of the deadlines.

CO-CHAIRPERSON MORAN: So while we're wrapping up, another thing is that almost everyone here is probably requesting some kind of reimbursement, and you got a brown manilla folder. And Linda Bunyan had provided you gory details about what it is you needed to supply, and even a mail-back envelope in there. So don't forget to do that. The sooner you do that, the sooner you'll get reimbursed.

It does take a little while. And the paperwork requirements are quite explicit, and you must do them exactly as they're -- including the blue ink signatures and all of this. So you'll get it back from Linda if you don't do that.

And it would help the Department. Scientists are the worst -- absolute worst at paperwork, and we all hate it, but the Department's been very kind to us in making the arrangements. I personally found it extremely generous and easy to be part of this meeting, and really appreciate the logistical work that the staff did in preparation here.

So one small thing we can do to help them finish that up is to do our forms properly and get them in quickly.

So that's it. Meg has a questions.

PANEL MEMBER SCHWARZMAN: I was just looking over the paperwork, and it says to send in receipts, and usually there's a cutoff below which you don't have to send in. Like the value of a receipt, like for the per diems for meals. Do you have to provide receipts?

MS. YEP: (Shakes head.)

PANEL MEMBER SCHWARZMAN: No. Thank you.

CO-CHAIRPERSON MORAN: Okay. So the answer was

25 | no, in case anyone didn't hear that.

All right. Are there other things that the staff wan to do before we close out the meeting here?

I can do that. I just wanted to know if you wanted to talk first or --

DEPUTY DIRECTOR WILLIAMS: No. Well, first of all, lunch -- I did want to mention lunch. So there's a table or two reserved at Blue Prynt Restaurant, if you'd like to join as group. And I'd kind of be curious to get a show of hands of folks who might be headed over to Blue Prynt. It's -- Blue Prynt is kitty corner. It's not -- it's kind of associated with Best Western, so it's right on that corner, if you happen to know where that is.

Are folks planning on going over there. We have a couple takers. So it will be probably a small-ish group, but yeah.

And I'd just -- this is my last chance I guess to say thank you, which is just tremendous. I'd heard great things about everybody on the Panel. And it certainly exceeded my already high expectations, in terms of what you might bring to the Department. And we are very, very fortunate to have all of you and your great thinking.

CO-CHAIRPERSON MORAN: And I also want to say thank you first to the panelists and then to the DTSC staff. And specifically, I'll start with logistical folks. They actually gave me a list to make sure that we

got everyone.

So Sharon, Heather -- Sharon Kipp, Heather
Kessler, Radhika Majhail, Jeff Wong, Linda Bunyan -- Jeff
is also doing science support here. Alexis Allston, Kim
Smith, and Corey Yep, in particular. Corey worked
exceptionally hard with us to get all this together.

And then I also really want to thank the professional staff. It is difficult to listen to a lot of people put in a critique of things you've done or are doing and to make statements that sound like you haven't thought about things, and we know you've thought about them and so on. And part of why I want to thank you so much is I know that there's a lot of work going forward. This is a long journey. And I want you to know -- I think I can say on behalf of all of us that we're coming on this journey with you. We're here to support you. We're going to be here to support you, and we really want to do that. So we're looking forward to that.

CO-CHAIRPERSON FONG: Yeah. I also want to thank the Panel members. I know how busy your schedules are and what you had to do in order to be here for a day and a half. And also for the people that actually had to travel across country. Flying is no fun these days. So I really appreciate you making the effort.

And in terms of the DTSC staff again, I just want

to mention the fact -- or reiterate what I said earlier. I was really amazed at how much work they put into organizing the agenda and the set up. It's impressive. And I'm never going to say that, you know, State workers done work again.

(Laughter.)

DEPUTY DIRECTOR WILLIAMS: I know you've never said that before.

PANEL MEMBER VERSTEEG: Never again.

PUBLIC PARTICIPATION SPECIALIST MAJHAIL: I have one more announcement to make for our listeners and for the members, that all the documents that shared today, all the PowerPoints, they will be on our website. So if they want -- you know, if you guys want to take a look later, they're going to be on the website.

Thank you.

PANEL MEMBER BLAKE: And because the co-chairs spent a lot of time thanking everybody else, I wanted to be sure that they got thanked as well. Thank you very much for taking on the work of being co-chair. I know it's no light lift here, so thank you for leading us through this next phase.

CO-CHAIRPERSON MORAN: Thank you very much. Is there any last thing anyone needs to say?

All RIGHT. This meeting is adjourned.

## CERTIFICATE OF REPORTER

I, JAMES F. PETERS, a Certified Shorthand
Reporter of the State of California, and Registered
Professional Reporter, do hereby certify:

That I am a disinterested person herein; that the foregoing California Department of Toxic Substance Control Green Ribbon Science Panel meeting was reported in shorthand by me, James F. Peters, a Certified Shorthand Reporter of the State of California, and thereafter transcribed under my direction, by computer-assisted transcription.

I further certify that I am not of counsel or attorney for any of the parties to said meeting nor in any way interested in the outcome of said meeting.

IN WITNESS WHEREOF, I have hereunto set my hand this 24th day of April, 2014.

James & Potter

JAMES F. PETERS, CSR, RPR
Certified Shorthand Reporter
License No. 10063